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(54) Title: COMPOSITIONS THAT INHIBIT PROLIFERATION OF CANCER CELLS

(57) Abstract: Disclosed are compositions and methods for reducing the proliferation of cancer cells through targeted interactions with integrins.



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COMPOSITIONS THAT INHIBIT PROLIFERATION OF CANCER CELLS

This application claims priority to United States Provisional Application No. 60/239,705 filed on October 12, 2000, entitled "Agents promoting apoptosis in cancer cells via interruption of oncogene-induced integrin signaling," which application is herein incorporated by reference in its entirety and to United States Provisional Application No. 60/242,812 filed on October 24, 2000 entitled, "Agents promoting apoptosis in cancer cells via interruption of oncogene-induced integrin signaling", and which application is herein incorporated by reference in its entirety.

I. BACKGROUND OF THE INVENTION

10 In cancer cells multiple oncogenic lesions cooperate in malignant transformation. Such cooperation permits survival and proliferation of tumor cells in absence of contact with extra-cellular matrix (ECM), suggesting that tumor cell survival and proliferation have become independent of the engagement of integrin signaling by ECM.

Carcinogenesis is caused by multiple cooperating genetic lesions leading to a progressive deregulation of cellular signaling and cell cycle restriction point control. The mutations involved result in oncogene activation or loss of tumor-suppressor gene function. Typically, single oncogenes are insufficient to cause malignant transformation because they simultaneously induce signals stimulating and inhibiting cell growth. As a result cell proliferation remains restricted. In contrast, cooperating oncogenic lesions act in concert to disable such inhibitory signals while reinforcing the growth-promoting stimuli. The co-operation of oncogenic lesions involves integration of multiple signals converging on the regulation of cell cycle-dependent kinase complexes (Lloyd *et al.*, 1997; Perez-Roger *et al.*, 1999; Roper *et al.*, 2001; Sewing *et al.*, 1997).

Disclosed herein are compositions and methods that show survival of various transformed cell types requires cell-autonomous (autocrine) integrin signaling activity. This activity is induced by cooperating oncogenic lesions and involves induction of integrin receptor and ligand components such as integrin alpha6 and integrin beta4, and laminin5-gamma2 chains. Blocking of integrin or the laminin ligand function induces rapid apoptosis of the transformed cells, even when growing in presence of ECM. In contrast, normal cells remain viable when exposed to the same treatment.

30 The disclosed compositions and methods are related to the cooperation of oncogenic lesions controlling the ability of transformed cells to proliferate in the absence of contact with the extra-cellular matrix (ECM). As taught herein, oncogenes cooperate to promote a cell-autonomous (autocrine) integrin signaling loop that proves essential for the survival of various transformed cell

types. As this signaling loop is not established in corresponding normal cells, the signaling components of this loop constitute attractive targets for cancer therapy.

II. SUMMARY OF THE INVENTION

In accordance with the purposes of this invention, as embodied and broadly described
5 herein, this invention, in one aspect, relates to compositions and methods related to integrin mediated cancer cell growth.

Additional advantages of the invention will be set forth in part in the description which follows, and in part will be obvious from the description, or may be learned by practice of the invention. The advantages of the invention will be realized and attained by means of the elements
10 and combinations particularly pointed out in the appended claims. It is to be understood that both the foregoing general description and the following detailed description are exemplary and explanatory only and are not restrictive of the invention, as claimed.

III. BRIEF DESCRIPTION OF THE DRAWINGS

The accompanying drawings, which are incorporated in and constitute a part of this
15 specification, illustrate several embodiments of the invention and together with the description, serve to explain the principles of the invention.

Figure 1 shows a series of schematics representative of the disclosed relationships and compositions. In normal cells integrin receptors signal to suppress programmed cell death (apoptosis) when engaged by appropriate extra-cellular matrix (ECM) ligands. When receptor-
20 ligand interaction is lost, the cells undergo apoptosis due to the lack of survival signals (Figure 1-. Panel 1). In cancer cells multiple oncogenic lesions cooperate to cause malignant transformation. Such cooperation permits survival and proliferation of tumor cells independent of integrin receptor-ECM interactions. This property has been termed anchorage-independence (Panel 2). We have discovered the mechanistic basis of anchorage-independence. The transformed cells
25 replace the requirement for ECM-dependent signaling with a surrogate integrin signaling loop on which they rely for survival. In colonic epithelial cells, activation of Ras in combination with APC (adenomatous polyposis coli) or p53 mutations leads to induction of integrin receptor and ligand components such as integrin alpha6 and laminin gamma2 chains. As a consequence, laminin-dependent activation of alpha6/beta4 integrin receptors signals to inhibit caspase activity and thus
30 to suppress apoptosis (Panel 3). Ablation of integrin alpha6, laminin ligand function or alpha6/beta4 integrin receptor function induces apoptosis of the transformed cells, even when growing in the presence of ECM. In contrast, non-transformed control cells remain viable when exposed to the same treatment, indicating that the dependence of the transformed cells on autocrine integrin signaling may be a particular feature of the cancer cell phenotype. The essential
35 role of alpha6 integrin extends to various transformed cell types including mesenchymal and

SW480 human colon carcinoma cells. Thus inhibition of laminin/alpha6 integrin-mediated signaling is an important method to induce cancer cell-specific death in a variety of cell types (Panel 4).

Figure 2 shows that oncogene cooperation protects cells from apoptosis. Figure 2A shows control, APCm, Ras, APCm+Ras cells that were detached from collagen IV substrates with Trypsin/EDTA and kept in suspension at 2×10^5 cells/ml in RPMI 10%FCS for 12 h at 39°C. Subsequently TUNEL analysis was performed on poly-lysine treated slides. The percentage of tunel-positive cells was determined by immunofluorescence microscopy. Figure 2B shows control, APCm, Ras, APCm Ras, SW480 cells were detached and maintained in suspension as described in (A). SW480 beta4 dn/Gal4VPER cells, express a 4OH-tamoxifen-inducible dominant-negative form of the beta4 integrin. Cells were pelleted and protein extracts were prepared in 300 μ l of 50 mM Tris-HCl, pH 7.4; 1% NP40 ; 0.25% sodium deoxycholate; 150 mM NaCl ; 1 mM EGTA. 200 μ l of extracts were incubated for 10 min with 2 μ l of Caspase 3 fluorometric Substrate (Upstate Biotechnology). Vmax of caspase activity was determined by measuring the fluorescence at 460 nm after excitation at 380 nm for 1 h. Figure 2C shows laminin gamma1 and gamma2-specific peptides were added to APCmin+Ras cells at the concentration of 100 μ g/ml. Caspase activity was measured as in (B). Figure 2D shows caspase 8 and caspase 9 activities were measured as in (B) using caspase 8 and caspase 9-specific fluorimetric substrates (Upstate Biotechnologies).

Figure 3 shows alterations of integrin and laminin expression profiles in malignant cell transformation. Figure 3A shows integrin expression and figure 3B shows laminin expression. The indicated cell populations were cultured on collagen IV-coated dishes at 39°C in RPMI 10% FCS. Total RNA was extracted from 10^6 cells for each sample and used for RT-PCR (laminins, alpha4, gamma2 and GAPDH) or RNase protection. For RT-PCR, cDNA was subjected to 28 cycles (linear range) of PCR amplification. PCR products were analyzed on a 2% agarose gel. For RNase protection, 10 μ g of total RNA was used per reaction. Products were resolved on a 4.5% polyacrylamide/10M urea gel.

Figure 4 shows alpha6/beta4 integrin is engaged by the laminin gamma2 chain to activate Shc (a src homology domain containing protein). Figure 4A shows cells detached from collagen IV coated dishes with 3mM EDTA (in PBS) were incubated on ice with the indicated antibodies for 1 h. Cells were then plated on 96well dishes coated with gamma1 or gamma2-specific peptides and were permitted to attach for 30 min in RPMI medium at 39°C. After incubation, wells were washed with RPMI. The percentage of attached cells per well was measured by hexoaminidase activity after lysis of the cells in the well and incubation with the substrate p-nitrophenol-N-acetyl- β -D-glucosaminide for 5 h at 37°C. Figure 4B shows cells indicated were detached as in (A) and

resuspended at 10^7 cells/ml in PBS. After 1 h of incubation on ice, cells were pelleted by centrifugation 5 min at 900 rpm and resuspended in RPMI containing the phosphatase inhibitor pervanadate at $10\mu\text{M}$. The cells were incubated with beta1 and beta4 integrin antibodies as well as the laminin gamma2-specific peptide for 40 min. Protein extracts were prepared and subjected to
 5 an immuno-precipitation with an anti-Shc antibody. Phospho-tyrosine was detected with a phospho-tyrosine-specific antibody in IP-western-blot (upper panels). The levels of Shc protein were monitored with a Shc-specific antibody reusing the same membrane (lower panels).

Figure 5 shows integrin alpha6/beta4 and Laminin gamma2 chain expression is essential for survival of transformed cells. Figure 5A1 shows the indicated cell populations were infected at
 10 an MOI of 2 with recombinant retroviruses expressing anti-sense RNA for alpha6 integrin, the gamma2 laminin chain or a beta4 integrin dominant-negative mutant together with a puromycin resistance gene. After 2 weeks of selection with puromycin, colonies were stained with Giemsa and counted. APCm+Ras alpha6 and APCm+Ras p35 cells express the alpha6 integrin chain and the anti-apoptotic baculovirus p35, respectively. APCm+Ras cells were also plated on dishes pre-
 15 coated with a laminin gamma2-specific peptide. Figure 5A2 shows APCm+Ras hygromycin, APCm+Ras Bcl2 and APCm+Ras p35 cells express the hygromycin resistance marker, and the anti-apoptotic proteins Bcl2 or p35, respectively. The cells were infected at an MOI of 2 with retroviruses carrying the beta4 integrin dominant-negative mutant and a puromycin resistance gene, or the puromycin resistance gene alone. Cells (10^5) were maintained in soft agar at 39°C for
 20 two weeks after which macroscopically visible colonies were counted. Figure 5A3 shows the indicated cell populations were infected at an MOI of 2 with recombinant retroviruses, drug-selected and maintained as described in A1. Figure 5A4 shows the indicated cell populations were infected at an MOI of 2 with recombinant retroviruses expressing anti-sense RNA for alpha6 integrin, the beta4 integrin dominant-negative mutant or the puromycin resistance marker. Cells
 25 (10^5) were maintained in soft agar and evaluated as described in A2. Figure 5B shows the effect of the alpha6 anti-sense RNA expression on the cell surface expression of alpha6 integrin was monitored by FACS analysis using an anti-alpha6 rat monoclonal antibody and a FITC conjugated goat anti-rat antiserum as the secondary antibody (upper panel). Expression of the beta4 dominant-negative mutant was confirmed by RNase protection. A probe overlapping the c-
 30 terminal end of the beta4 dominant negative mRNA was used to measure beta4 integrin and beta4 dn expression in the same sample (lower panel). Figure 5C shows arf null mouse embryo fibroblasts (MEFs) that were infected with retroviruses expressing the oncogenes Myc and Ras. Cells were then additionally infected with a retroviruses expressing anti-sense RNA for alpha6 integrin or the beta4 integrin dominant-negative mutant as shown in (A). MEFs
 35 Ras/Myc/Arfnull+alpha6 sense express the alpha6 integrin chain. Figure 4D shows SW480 human

colon carcinoma cells that were infected as described in (A) using VSV pseudo-typed viruses. Infected cells were selected with puromycin in soft agar. Clones were counted 2 weeks after selection. SW480 beta4 dn/Gal4VPER cells, express a 4OH-tamoxifen-inducible dominant-negative form of the beta4 integrin. The cells (10^5) were maintained in soft agar for 2 weeks in
5 presence or absence of 4OH-tamoxifen.

IV. DETAILED DESCRIPTION

The present invention may be understood more readily by reference to the following detailed description of preferred embodiments of the invention and the Examples included therein and to the Figures and their previous and following description.

10 Before the present compounds, compositions, articles, devices, and/or methods are disclosed and described, it is to be understood that this invention is not limited to specific synthetic methods, specific recombinant biotechnology methods unless otherwise specified, or to particular reagents unless otherwise specified, as such may, of course, vary. It is also to be understood that the terminology used herein is for the purpose of describing particular embodiments only and is
15 not intended to be limiting.

A. Definitions

As used in the specification and the appended claims, the singular forms "a," "an" and "the" include plural referents unless the context clearly dictates otherwise. Thus, for example, reference to "a pharmaceutical carrier" includes mixtures of two or more such carriers, and the
20 like.

Ranges may be expressed herein as from "about" one particular value, and/or to "about" another particular value. When such a range is expressed, another embodiment includes from the one particular value and/or to the other particular value. Similarly, when values are expressed as approximations, by use of the antecedent "about," it will be understood that the particular value
25 forms another embodiment. It will be further understood that the endpoints of each of the ranges are significant both in relation to the other endpoint, and independently of the other endpoint.

In this specification and in the claims which follow, reference will be made to a number of terms which shall be defined to have the following meanings:

"Optional" or "optionally" means that the subsequently described event or circumstance
30 may or may not occur, and that the description includes instances where said event or circumstance occurs and instances where it does not.

"Primers" are a subset of probes which are capable of supporting some type of enzymatic manipulation and which can hybridize with a target nucleic acid such that the enzymatic manipulation can occur. A primer can be made from any combination of nucleotides or nucleotide

derivatives or analogs available in the art which do not interfere with the enzymatic manipulation.

“Probes” are molecules capable of interacting with a target nucleic acid, typically in a sequence specific manner, for example through hybridization. The hybridization of nucleic acids is well understood in the art and discussed herein. Typically a probe can be made from any
5 combination of nucleotides or nucleotide derivatives or analogs available in the art.

B. Compositions and methods

Disclosed are compositions and methods related to integrins and integrin signaling. It is shown herein that integrin alpha6, integrin beta4, and laminin5, through at least up regulation of the beta and gamma chains of laminin5, are upregulated in cancer cells. Integrin alpha6 and
10 integrin beta4 interact to form the integrin receptor, A6B4. The integrin receptor A6B4 specifically interacts with laminin5 and laminin5 specifically interacts with A6B4. Furthermore, it is shown that interference with the production or function of alpha6, beta4, or the laminin gamma2 chain not only prevents proliferation of the cancer cells, dependent on the upregulation of alpha6, beta4, and the laminin gamma2 chain, but that this kills the cancer cells as well. Alpha6 and beta4
15 signaling occur through the integrin receptor A6B4. Thus, interference with the formation of A6B4, will interfere with the function of A6B4, for example, the signaling of A6B4. Thus, disclosed are compositions and methods that interfere with the function of alpha6, beta4, laminin5, the laminin gamma2 chain, or A6B4. Also disclosed are compositions and methods that interfere with the function of molecules involved with the signal transduction that is connected to either
20 alpha6, beta4, laminin5, the laminin gamma2 chain, or A6B4. Also disclosed are methods for reducing the proliferation of cancer cells, as well as methods of killing cancer cells that involve using the compositions disclosed herein that interfere, reduce, or eliminate the function or the alpha6, beta4, laminin5, the laminin gamma2 chain, or A6B4 function.

Disclosed herein is a relationship between two types of molecules in a cancer cell. The
25 first type of molecule is an integrin receptor, composed of integrins, and the second type of molecule is a ligand that interacts with the integrin receptor, through the integrins. There are specificities that exist between the integrin receptors and their ligands. One aspect, disclosed herein is that when a cell goes from a non-cancerous state to a cancerous state, there is a co-upregulation of both the ligand (or parts of the ligand, such as subunits) and the cognate integrin
30 receptor. The co-upregulation of both types of molecules creates an autocrine loop situation, wherein the signaling pathways controlled by the integrin receptor become autonomously activated, rather than exogenously activated, as would normally occur. The upregulation of both types of molecules creates a more fully transformed cellular phenotype in which cancer cell survival depends on the autocrine loop. Now therapeutic activities can target both points in the
35 autocrine loop. Specific examples, of this co-upregulation in cancer cells are disclosed herein. For

example, laminin 5 (both the beta and gamma2 chains are upregulated) and integrin receptor alpha6beta4; laminin 10/11 and the integrin receptors alpha6beta1 and alpha3beta1.

As the Examples herein indicate which integrins, which integrin receptor, and which integrin ligand are involved in conferring cancer cells ability to grow in the absence of extra cellular matrix (ECM), also disclosed are methods using these integrins, integrin receptors and ligands to identify molecules that interact with them and/or interfere with their function.

1. Compositions

Disclosed are the components to be used to prepare the disclosed compositions as well as the compositions themselves to be used within the methods disclosed herein. These and other materials are disclosed herein, and it is understood that when combinations, subsets, interactions, groups, etc. of these materials are disclosed, while specific reference of each various individual and collective permutation of these compounds may not be explicitly disclosed, each is specifically contemplated and described herein. For example, if a particular beta4 or alpha6 is disclosed and discussed and a number of modifications that can be made to a number of molecules including the modifications to beta4 or alpha6 are discussed, specifically contemplated is each and every combination and permutation of these modifications and the modifications that are possible unless specifically indicated to the contrary. Thus, if a class of molecules A, B, and C are disclosed as well as a class of molecules D, E, and F and an example of a combination molecule, A-D is disclosed, then even if each is not individually recited, each is individually and collectively contemplated. Thus, combinations, A-E, A-F, B-D, B-E, B-F, C-D, C-E, and C-F are considered disclosed. Likewise, any subset or combination of these is also disclosed. Thus, for example, the sub-group of A-E, B-F, and C-E would be considered disclosed. This concept applies to all aspects of this application including, but not limited to, steps in methods of making and using the disclosed compositions. Thus, if there are a variety of additional steps that can be performed it is understood that each of these additional steps can be performed with any specific embodiment or combination of embodiments of the disclosed methods.

a) Integrins and their ligands

For proper embryonic development, tissue homeostasis, and wound healing, cell proliferation must be tightly regulated, both in space and over time. In particular, a cell must be able to sense its relationship to other cells and the extra-cellular matrix (ECM) and convert these positional cues into biochemical signals affecting the regulation of proliferation. Because of their ability to couple the recognition of positional cues to the activation of intracellular signaling pathways, adhesion receptors, such as integrins and cadherins, are likely to be necessary to achieve this goal.

The integrins mediate cell adhesion primarily by binding to distinct, although overlapping,

subsets of ECM proteins. Normal cells require contact with serum-derived ECM components for proliferation, differentiation and survival (e.g. Clark and Brugge, 1995; Lin and Bissell, 1993; Parise *et al.*, 2000), a phenomenon called anchorage-dependence. This involves signaling through integrin receptors (Hynes, 1992). Fibronectin and laminin as well as other ECM proteins are
5 known to act as ligands for integrin receptors (Akiyama *et al.*, 1990). Integrins are transmembrane proteins forming alpha-beta chain heterodimers. Alpha and beta chain integrins are members of distinct gene families. The ligand binding specificity of the hetero dimers is determined by specific combinations of alpha and beta chain gene family members (Hynes, 1992). Ligand binding triggers signaling of integrin receptors through the cytoplasmic tail of the beta chain via
10 interaction with various signaling components.

Integrins activate common as well as subgroup-specific signaling pathways (Clark and Brugge, 1995; Giancotti and Ruoslahti, 1999). In particular, while most integrins activate focal adhesion kinase (FAK), the $\alpha 1\beta 1$, $\alpha 5\beta 1$, $\alpha v\beta 3$ and $\alpha 6\beta 4$ integrins are coupled to the Ras-extracellular signal-regulated kinase (ERK) signaling pathway by Shc (Mainiero *et al.*, 1997;
15 Mainiero *et al.*, 1995; Wary *et al.*, 1996). Shc is an SH2-PTB domain adapter protein expressed in three forms, p46, p52 and p66, two of which (p46 and p52) link various tyrosine kinases to Ras by recruiting the Grb2/SOS complex to the plasma membrane (Pawson and Scott, 1997). Upon activation by SOS, Ras stimulates a kinase cascade culminating in the activation of the mitogen-activated protein kinase (MAPK) ERK (Marshall, 1995). ERK phosphorylates ternary complex
20 transcription factors, such as Elk-1 and Sap-1/2, and promotes transcription of the immediate-early gene Fos (Treisman, 1996). In primary endothelial cells and keratinocytes, mitogens and Shc-linked integrins cooperate, in a synergic fashion, to promote transcription from the Fos promoter. Accordingly, ligation of integrins linked to Shc enables these cells to progress through G1 in response to mitogens, whereas ligation of other integrins results in growth arrest, even in the
25 presence of mitogens (Mainiero *et al.*, 1997; Wary *et al.*, 1996). Shc is like a binary switch controlling cell cycle progression in response to the ECM. Moreover, integrin receptors have been shown to induce intracellular signaling leading to AKT activation supporting cell survival (Lee and Juliano, 2000).

In contrast with normal cells, cancer cells generally are able to survive and proliferate in
30 the absence of anchorage to ECM (Giancotti and Mainiero, 1994), suggesting that tumor cell survival and proliferation have become independent of the engagement of integrin signaling through ECM.

Proliferation in the absence of anchorage to ECM of secondary rat embryo fibroblasts requires the cooperation of Ras and Myc or Ras and adenovirus E1a oncogenes (Land *et al.*, 1983;
35 Ruley, 1983). Similarly, murine colonic epithelial cells require both activated Ras and mutation of

the adenomous polyposis coli gene (APC^{min}) (D'Abaco et al., 1996) in order to proliferate in suspension.

Integrins are a large family of cell surface receptor molecules that function to mediate interactions between cells and between cells and the extracellular matrix. Integrin receptors are
5 heterodimers composed of two subunits, an alpha integrin and a beta integrin. The heterodimer forms, is expressed on the cell surface, and acts to transmit signals obtained from interactions with the extracellular matrix or other cells, through the cellular membrane and into the cytosol of the cell. The signal transduction that takes place occurs because of ligand interactions with the receptor. Integrin receptors can have a number of ligands, including collagens, fibronectins, and
10 laminins.

There are currently at least 18 different alpha integrins and at least 8 different beta integrins that have been shown to form at least 24 different alphabeta heterodimers. Certain integrins, such as beta1, interact in a number of different heterodimers, but many subunits only form a single heterodimer, either because of structural constraints on their interactions, or cellular
15 expression patterns that provide only a limited number of potential dimer partners. Disclosed herein are specific relationships that occur between a subset of integrins, integrin receptors, and their ligands. The disclosed relationships, revolve around the alpha6beta4 receptor, formed by the alpha6 and beta4 integrins. Of particular interest is the relationship between the ligand for the alpha6beta4 receptor, laminin5. The laminins are made up of 3 chains, an alpha chain, a beta
20 chain, and a gamma chain. The specificity of the interaction between laminin5 and alpha6beta4 receptor is controlled by the gamma chain. Laminin5 contains a gamma2 chain which only interacts with the alpha6beta4 integrin receptor.

Integrin alpha6 has seven amino-terminal repeating segments that may fold into a seven unit beta-propeller, five n-terminal FG-GAP domains and three divalent cation sites. The
25 transmembrane domain is followed by a short cytoplasmic tail, that is alternatively spliced in A and B forms. The alpha6 integrin chain also shows alternative splicing between repeat units III and IV, resulting in the presence or absence of Exon X2. Integrin alpha6 is processed into a heavy and a light chain that are disulphide linked. A representative allele of the human alpha6 cDNA is set forth in SEQ ID NO:1. It is understood that the disclosed functional domains as well as the
30 others contained within alpha6 are considered separately disclosed as discrete fragments of the alpha6 protein as well as the nucleic acid that encodes them.

Integrin beta4 contains a MIDAS-like motif and four cysteine-rich repeats, three EGF-like domains in the N-terminal extracellular domain, a trans-membrane region and a long cytoplasmic tail containing two pairs of fibronectin Type III repeats. The latter are connected by a variable
35 segment that may undergo alternative splicing. Integrin beta4 also undergoes proteolytic

processing in its cytoplasmic tail, causing the 200kD mature form to be converted to 165 and 130kD fragments. A representative allele of the human beta4 cDNA is set forth in SEQ ID NO:5. It is understood that the disclosed functional domains as well as the others contained within beta4 are considered separately disclosed as discreet fragments of the beta4 protein as well as the nucleic acid that encodes them.

Laminin5 is composed of the laminin chains alpha3, beta3 and gamma2. Laminin5 can may contain either the shorter laminin alpha3A chain or the longer alpha3B chain. Laminin5 can also be trimmed by proteolytic processing of the N-terminal portion of its alpha3A chain and the N-terminal portion of the gamma2 chain. The Laminin gamma2 chain contains at least six laminin EGF-like domains (Domains III and V) with an embedded laminin B domain (Domain IV) within the N-terminal half. The c-terminal tail contains a coiled-coil domain. The N-terminal processed portion of the gamma2 chain is sufficient to bind to and activate the integrin alpha6/beta4 receptor. A representative allele of the human laminin5-gamma2 cDNA is set forth in SEQ ID NO:13. It is understood that the disclosed functional domains as well as the others contained within the laminin5 protein and laminin5-gamma2 protein are considered separately disclosed as discreet fragments of the laminin5 protein and laminin5-gamma2 protein as well as and the nucleic acid that encodes them.

Disclosed are compositions and methods for inhibiting integrin signaling, for example, integrin signaling dependent on alpha6 and beta4 integrins. For example, compositions and methods that inhibit integrin receptor signaling from, for example, the alpha6beta4 integrin receptor are disclosed. It is also understood that the integrin receptor signaling can be affected by, for example, interfering with a molecule, such as a ligand for the integrin receptor, or a downstream signaling molecule of the integrin receptor in a way that prevents the integrin receptor signal from being fully propagated. It is understood that the compositions and methods for inhibition of integrin signaling and function can be any composition or method that ultimately inhibits the cell proliferation in which the integrin is expressed, by for example killing the cell. It is understood that the compositions and methods typically can fall into three basic non-limiting classes of function regulators, which are discussed herein.

(1) Classes

(a) Production regulators

Production regulators is a broad class of integrin function regulators that are directed at the production of the target integrin, by for example, preventing mRNA synthesis or expression of the target integrin, or by causing mRNA degradation of the target integrin which inhibits the translation of the target integrin. While production regulators, can be any type of molecule targeting any point in the integrin production pathway, typically these types of compositions will

target either the mRNA expression or the protein expression of the integrin. For example, if beta4 integrin, alpha6 integrin, or the gamma2 subunit of laminin5, which has been shown herein to be upregulated in cancer cells and which causes the cancer cell to be able to live in the absence of the ECM, was the target integrin, a typical production regulator of beta4 integrin, alpha6 integrin, or the gamma2 subunit of laminin5 would be for example, an antisense molecule that targeted the mRNA of beta4 integrin, alpha6 integrin, or the gamma2 subunit of laminin5. It is also understood that a production regulator could also target any molecule that is disclosed herein, or is within the signaling chain associated with a target integrin or integrin receptor. For example, the inhibition of the production of the ligand for a target integrin receptor is one way of inhibiting integrin function of that receptor. Thus, production regulators can either inhibit or enhance integrin production.

(b) Integrin to integrin regulators

Another type of integrin function regulator is an integrin-integrin regulator. This type of function regulator, typically prevents integrins from interacting to form a functional integrin receptor. For example, an integrin to integrin regulator could be a composition that would interact with beta4 in a way that would prevent beta4 from interacting with alpha6 to form the alpha6beta4 integrin receptor or it could be a composition that would interact with alpha6 in a way that would prevent alpha6 from interacting with beta4 to form the alpha6beta4 integrin receptor. It is also contemplated that the function regulators of the integrin to integrin interaction can affect the signaling pathways dependent on integrin receptors containing alpha6 or beta4 integrins. It is not required that an integrin to integrin regulator prevent an integrin from interacting with all of the possible integrin partners it could interact with, just that it prevent the interaction of the target integrin with another specific integrin. For example, the integrin alpha6 interacts with beta1 integrin and beta4 integrin. In certain embodiments, the compositions interfere with alpha6-beta4 interactions but do not interfere with alpha6-beta1 interactions.

(c) Integrin to other molecule regulators

The third class of function regulators are the integrin to other molecule regulators. These compositions are designed to specifically interfere with molecules such as small molecule ligands or other proteins that interact with the integrin or integrin receptor. For example, an integrin to other molecule regulator might target a ligand for a particular integrin receptor, such as the alpha6beta4 receptor. The ligand for the alpha6beta4 receptor is laminin5 or the laminin gamma2 chain. Compositions that interact with laminin5 such that laminin5 or the laminin gamma2 chain interactions with alpha6beta4 are inhibited or reduced are specifically contemplated herein. Likewise, there are other molecules, such as Shc molecules, that also interact with integrins, such as the alpha6beta4 receptor. Compositions that specifically interact with the Shc molecules such that they prevent the appropriate interactions between the Shc molecule and the alpha6beta4

receptor are disclosed.

(2) Types

Just as there are different general classes of molecules that can regulate (such as by inhibition) the function of the disclosed integrins and integrin receptors, so to there are many different types of molecules that perform that regulation. For example, any molecule that can perform the regulation of for example, the disclosed integrins, integrin receptors, or signaling pathways produced by the disclosed integrins and integrin receptors are contemplated. For example, antibodies or small molecules which inhibit the disclosed compositions are herein disclosed. Also disclosed are, for example, functional nucleic acids, such as ribozymes or antisense molecules that can inhibit the disclosed integrin function in a variety of ways. A non-limiting list of exemplary molecules is discussed herein.

(a) Antibodies

Antibodies can be used to regulate, for example, the function of the disclosed integrins and integrin receptors, molecules that interact with the disclosed integrin receptors, and molecules in the signaling pathways of the disclosed integrin receptors.

As used herein, the term "antibody" encompasses, but is not limited to, whole immunoglobulin (i.e., an intact antibody) of any class. Native antibodies are usually heterotetrameric glycoproteins, composed of two identical light (L) chains and two identical heavy (H) chains. Typically, each light chain is linked to a heavy chain by one covalent disulfide bond, while the number of disulfide linkages varies between the heavy chains of different immunoglobulin isotypes. Each heavy and light chain also has regularly spaced intrachain disulfide bridges. Each heavy chain has at one end a variable domain (V(H)) followed by a number of constant domains. Each light chain has a variable domain at one end (V(L)) and a constant domain at its other end; the constant domain of the light chain is aligned with the first constant domain of the heavy chain, and the light chain variable domain is aligned with the variable domain of the heavy chain. Particular amino acid residues are believed to form an interface between the light and heavy chain variable domains. The light chains of antibodies from any vertebrate species can be assigned to one of two clearly distinct types, called kappa (k) and lambda (l), based on the amino acid sequences of their constant domains. Depending on the amino acid sequence of the constant domain of their heavy chains, immunoglobulins can be assigned to different classes. There are five major classes of human immunoglobulins: IgA, IgD, IgE, IgG and IgM, and several of these may be further divided into subclasses (isotypes), e.g., IgG-1, IgG-2, IgG-3, and IgG-4; IgA-1 and IgA-2. One skilled in the art would recognize the comparable classes for mouse. The heavy chain constant domains that correspond to the different classes of immunoglobulins are called alpha, delta, epsilon, gamma, and mu, respectively.

Antibodies can be either polyclonal or monoclonal. Polyclonal antibodies, typically are derived from the serum of an animal that has been immunogenically challenged, and monoclonal antibodies are derived as discussed herein.

The term "variable" is used herein to describe certain portions of the variable domains that
5 differ in sequence among antibodies and are used in the binding and specificity of each particular antibody for its particular antigen. However, the variability is not usually evenly distributed through the variable domains of antibodies. It is typically concentrated in three segments called complementarity determining regions (CDRs) or hypervariable regions both in the light chain and the heavy chain variable domains. The more highly conserved portions of the variable domains
10 are called the framework (FR). The variable domains of native heavy and light chains each comprise four FR regions, largely adopting a b-sheet configuration, connected by three CDRs, which form loops connecting, and in some cases forming part of, the b-sheet structure. The CDRs in each chain are held together in close proximity by the FR regions and, with the CDRs from the other chain, contribute to the formation of the antigen binding site of antibodies (see Kabat E. A.
15 et al., "Sequences of Proteins of Immunological Interest," National Institutes of Health, Bethesda, Md. (1987)). The constant domains are not involved directly in binding an antibody to an antigen, but exhibit various effector functions, such as participation of the antibody in antibody-dependent cellular toxicity.

As used herein, the term "antibody or fragments thereof" encompasses chimeric antibodies
20 and hybrid antibodies, with dual or multiple antigen or epitope specificities, and fragments, such as F(ab')₂, Fab', Fab and the like, including hybrid fragments. Thus, fragments of the antibodies that retain the ability to bind their specific antigens are provided. For example, fragments of antibodies which maintain EphA2 binding activity are included within the meaning of the term "antibody or fragment thereof." Such antibodies and fragments can be made by techniques known in the art and
25 can be screened for specificity and activity according to the methods set forth in the Examples and in general methods for producing antibodies and screening antibodies for specificity and activity (See Harlow and Lane. Antibodies, A Laboratory Manual. Cold Spring Harbor Publications, New York, (1988)).

Also included within the meaning of "antibody or fragments thereof" are conjugates of
30 antibody fragments and antigen binding proteins (single chain antibodies) as described, for example, in U.S. Pat. No. 4,704,692, the contents of which are hereby incorporated by reference. Single chain divalent antibodies are also provided.

Optionally, the antibodies are generated in other species and "humanized" for
administration in humans. Humanized forms of non-human (e.g., murine) antibodies are chimeric
35 immunoglobulins, immunoglobulin chains or fragments thereof (such as Fv, Fab, Fab', F(ab')₂, or

other antigen-binding subsequences of antibodies) which contain minimal sequence derived from non-human immunoglobulin. Humanized antibodies include human immunoglobulins (recipient antibody) in which residues from a complementary determining region (CDR) of the recipient are replaced by residues from a CDR of a non-human species (donor antibody) such as mouse, rat or
5 rabbit having the desired specificity, affinity and capacity. In some instances, Fv framework residues of the human immunoglobulin are replaced by corresponding non-human residues. Humanized antibodies may also comprise residues that are found neither in the recipient antibody nor in the imported CDR or framework sequences. In general, the humanized antibody will comprise substantially all of at least one, and typically two, variable domains, in which all or
10 substantially all of the CDR regions correspond to those of a non-human immunoglobulin and all or substantially all of the FR regions are those of a human immunoglobulin consensus sequence. The humanized antibody optimally also will comprise at least a portion of an immunoglobulin constant region (Fc), typically that of a human immunoglobulin (Jones et al., *Nature*, 321:522-525 (1986); Riechmann et al., *Nature*, 332:323-327 (1988); and Presta, *Curr. Op. Struct. Biol.*, 2:593-
15 596 (1992)).

Methods for humanizing non-human antibodies are well known in the art. Generally, a humanized antibody has one or more amino acid residues introduced into it from a source that is non-human. These non-human amino acid residues are often referred to as "import" residues, which are typically taken from an "import" variable domain. Humanization can be essentially
20 performed following the method of Winter and co-workers (Jones et al., *Nature*, 321:522-525 (1986); Riechmann et al., *Nature*, 332:323-327 (1988); Verhoeyen et al., *Science*, 239:1534-1536 (1988)), by substituting rodent CDRs or CDR sequences for the corresponding sequences of a human antibody. Accordingly, such "humanized" antibodies are chimeric antibodies (U.S. Pat. No. 4,816,567), wherein substantially less than an intact human variable domain has been
25 substituted by the corresponding sequence from a non-human species. In practice, humanized antibodies are typically human antibodies in which some CDR residues and possibly some FR residues are substituted by residues from analogous sites in rodent antibodies.

The choice of human variable domains, both light and heavy, to be used in making the humanized antibodies is very important in order to reduce antigenicity. According to the "best-fit"
30 method, the sequence of the variable domain of a rodent antibody is screened against the entire library of known human variable domain sequences. The human sequence which is closest to that of the rodent is then accepted as the human framework (FR) for the humanized antibody (Sims et al., *J. Immunol.*, 151:2296 (1993) and Chothia et al., *J. Mol. Biol.*, 196:901 (1987)). Another method uses a particular framework derived from the consensus sequence of all human antibodies
35 of a particular subgroup of light or heavy chains. The same framework may be used for several different humanized antibodies (Carter et al., *Proc. Natl. Acad. Sci. USA*, 89:4285 (1992);

Presta et al., J. Immunol., 151:2623 (1993)).

It is further important that antibodies be humanized with retention of high affinity for the antigen and other favorable biological properties. To achieve this goal, according to a preferred method, humanized antibodies are prepared by a process of analysis of the parental sequences and
5 various conceptual humanized products using three dimensional models of the parental and humanized sequences. Three dimensional immunoglobulin models are commonly available and are familiar to those skilled in the art. Computer programs are available which illustrate and display probable three-dimensional conformational structures of selected candidate immunoglobulin sequences. Inspection of these displays permits analysis of the likely role of the
10 residues in the functioning of the candidate immunoglobulin sequence, i.e., the analysis of residues that influence the ability of the candidate immunoglobulin to bind its antigen. In this way, FR residues can be selected and combined from the consensus and import sequence so that the desired antibody characteristic, such as increased affinity for the target antigen(s), is achieved. In general, the CDR residues are directly and most substantially involved in influencing antigen binding (see,
15 WO 94/04679, published 3 March 1994).

Transgenic animals (e.g., mice) that are capable, upon immunization, of producing a full repertoire of human antibodies in the absence of endogenous immunoglobulin production can be employed. For example, it has been described that the homozygous deletion of the antibody heavy chain joining region (J(H)) gene in chimeric and germ-line mutant mice results in complete
20 inhibition of endogenous antibody production. Transfer of the human germ-line immunoglobulin gene array in such germ-line mutant mice will result in the production of human antibodies upon antigen challenge (see, e.g., Jakobovits et al., Proc. Natl. Acad. Sci. USA, 90:2551-255 (1993); Jakobovits et al., Nature, 362:255-258 (1993); Bruggemann et al., Year in Immuno., 7:33 (1993)). Human antibodies can also be produced in phage display libraries (Hoogenboom et al., J. Mol.
25 Biol., 227:381 (1991); Marks et al., J. Mol. Biol., 222:581 (1991)). The techniques of Cote et al. and Boerner et al. are also available for the preparation of human monoclonal antibodies (Cole et al., Monoclonal Antibodies and Cancer Therapy, Alan R. Liss, p. 77 (1985); Boerner et al., J. Immunol., 147(1):86-95 (1991)).

The present invention further provides a hybridoma cell that produces the monoclonal
30 antibody of the invention. The term "monoclonal antibody" as used herein refers to an antibody obtained from a substantially homogeneous population of antibodies, i.e., the individual antibodies comprising the population are identical except for possible naturally occurring mutations that may be present in minor amounts. The monoclonal antibodies herein specifically include "chimeric" antibodies in which a portion of the heavy and/or light chain is identical with or homologous to
35 corresponding sequences in antibodies derived from a particular species or belonging to a

particular antibody class or subclass, while the remainder of the chain(s) is identical with or homologous to corresponding sequences in antibodies derived from another species or belonging to another antibody class or subclass, as well as fragments of such antibodies, so long as they exhibit the desired activity (See, U.S. Pat. No. 4,816,567 and Morrison et al., Proc. Natl. Acad. Sci. USA, 81:6851-6855 (1984)).

Monoclonal antibodies of the invention may be prepared using hybridoma methods, such as those described by Kohler and Milstein, Nature, 256:495 (1975) or Harlow and Lane. Antibodies, A Laboratory Manual. Cold Spring Harbor Publications, New York, (1988). In a hybridoma method, a mouse or other appropriate host animal, is typically immunized with an immunizing agent to elicit lymphocytes that produce or are capable of producing antibodies that will specifically bind to the immunizing agent. Alternatively, the lymphocytes may be immunized in vitro. Preferably, the immunizing agent comprises EphA2. Traditionally, the generation of monoclonal antibodies has depended on the availability of purified protein or peptides for use as the immunogen. More recently DNA based immunizations have shown promise as a way to elicit strong immune responses and generate monoclonal antibodies. In this approach, DNA-based immunization can be used, wherein DNA encoding a portion of EphA2 expressed as a fusion protein with human IgG1 is injected into the host animal according to methods known in the art (e.g., Kilpatrick KE, et al. Gene gun delivered DNA-based immunizations mediate rapid production of murine monoclonal antibodies to the Flt-3 receptor. Hybridoma. 1998 Dec;17(6):569-76; Kilpatrick KE et al. High-affinity monoclonal antibodies to PED/PEA-15 generated using 5 microg of DNA. Hybridoma. 2000 Aug;19(4):297-302, which are incorporated herein by referenced in full for the methods of antibody production).

An alternate approach to immunizations with either purified protein or DNA is to use antigen expressed in baculovirus. The advantages to this system include ease of generation, high levels of expression, and post-translational modifications that are highly similar to those seen in mammalian systems. Use of this system involves expressing domains of EphA2 antibody as fusion proteins. The antigen is produced by inserting a gene fragment in-frame between the signal sequence and the mature protein domain of the EphA2 antibody nucleotide sequence. This results in the display of the foreign proteins on the surface of the virion. This method allows immunization with whole virus, eliminating the need for purification of target antigens.

Generally, peripheral blood lymphocytes ("PBLs") are used in methods of producing monoclonal antibodies if cells of human origin are desired, or spleen cells or lymph node cells are used if non-human mammalian sources are desired. The lymphocytes are then fused with an immortalized cell line using a suitable fusing agent, such as polyethylene glycol, to form a hybridoma cell (Goding, "Monoclonal Antibodies: Principles and Practice" Academic Press,

(1986) pp. 59-103). Immortalized cell lines are usually transformed mammalian cells, including myeloma cells of rodent, bovine, equine, and human origin. Usually, rat or mouse myeloma cell lines are employed. The hybridoma cells may be cultured in a suitable culture medium that preferably contains one or more substances that inhibit the growth or survival of the unfused, 5 immortalized cells. For example, if the parental cells lack the enzyme hypoxanthine guanine phosphoribosyl transferase (HGPRT or HPRT), the culture medium for the hybridomas typically will include hypoxanthine, aminopterin, and thymidine ("HAT medium"), which substances prevent the growth of HGPRT-deficient cells. Preferred immortalized cell lines are those that fuse efficiently, support stable high level expression of antibody by the selected antibody-producing 10 cells, and are sensitive to a medium such as HAT medium. More preferred immortalized cell lines are murine myeloma lines, which can be obtained, for instance, from the Salk Institute Cell Distribution Center, San Diego, Calif. and the American Type Culture Collection, Rockville, Md. Human myeloma and mouse-human heteromyeloma cell lines also have been described for the production of human monoclonal antibodies (Kozbor, J. Immunol., 133:3001 (1984); Brodeur et al., "Monoclonal Antibody Production Techniques and Applications" Marcel Dekker, Inc., New 15 York, (1987) pp. 51-63).

The culture medium in which the hybridoma cells are cultured can then be assayed for the presence of monoclonal antibodies directed against EphA2. Preferably, the binding specificity of monoclonal antibodies produced by the hybridoma cells is determined by immunoprecipitation or 20 by an in vitro binding assay, such as radioimmunoassay (RIA) or enzyme-linked immunoabsorbent assay (ELISA). Such techniques and assays are known in the art, and are described further in the Examples below or in Harlow and Lane "Antibodies, A Laboratory Manual" Cold Spring Harbor Publications, New York, (1988).

After the desired hybridoma cells are identified, the clones may be subcloned by limiting 25 dilution or FACS sorting procedures and grown by standard methods. Suitable culture media for this purpose include, for example, Dulbecco's Modified Eagle's Medium and RPMI-1640 medium. Alternatively, the hybridoma cells may be grown in vivo as ascites in a mammal.

The monoclonal antibodies secreted by the subclones may be isolated or purified from the culture medium or ascites fluid by conventional immunoglobulin purification procedures such as, 30 for example, protein A-Sepharose, protein G, hydroxylapatite chromatography, gel electrophoresis, dialysis, or affinity chromatography.

The monoclonal antibodies may also be made by recombinant DNA methods, such as those described in U.S. Pat. No. 4,816,567. DNA encoding the monoclonal antibodies of the invention can be readily isolated and sequenced using conventional procedures (e.g., by using 35 oligonucleotide probes that are capable of binding specifically to genes encoding the heavy and

light chains of murine antibodies). The hybridoma cells of the invention serve as a preferred source of such DNA. Once isolated, the DNA may be placed into expression vectors, which are then transfected into host cells such as simian COS cells, Chinese hamster ovary (CHO) cells, plasmacytoma cells, or myeloma cells that do not otherwise produce immunoglobulin protein, to obtain the synthesis of monoclonal antibodies in the recombinant host cells. The DNA also may be modified, for example, by substituting the coding sequence for human heavy and light chain constant domains in place of the homologous murine sequences (U.S. Pat. No. 4,816,567) or by covalently joining to the immunoglobulin coding sequence all or part of the coding sequence for a non-immunoglobulin polypeptide. Optionally, such a non-immunoglobulin polypeptide is substituted for the constant domains of an antibody of the invention or substituted for the variable domains of one antigen-combining site of an antibody of the invention to create a chimeric bivalent antibody comprising one antigen-combining site having specificity for EphA2 and another antigen-combining site having specificity for a different antigen.

In vitro methods are also suitable for preparing monovalent antibodies. Digestion of antibodies to produce fragments thereof, particularly, Fab fragments, can be accomplished using routine techniques known in the art. For instance, digestion can be performed using papain. Examples of papain digestion are described in WO 94/29348 published Dec. 22, 1994, U.S. Pat. No. 4,342,566, and Harlow and Lane, *Antibodies, A Laboratory Manual*, Cold Spring Harbor Publications, New York, (1988). Papain digestion of antibodies typically produces two identical antigen binding fragments, called Fab fragments, each with a single antigen binding site, and a residual Fc fragment. Pepsin treatment yields a fragment, called the F(ab')₂ fragment, that has two antigen combining sites and is still capable of cross-linking antigen.

The Fab fragments produced in the antibody digestion also contain the constant domains of the light chain and the first constant domain of the heavy chain. Fab' fragments differ from Fab fragments by the addition of a few residues at the carboxy terminus of the heavy chain domain including one or more cysteines from the antibody hinge region. The F(ab')₂ fragment is a bivalent fragment comprising two Fab' fragments linked by a disulfide bridge at the hinge region. Fab'-SH is the designation herein for Fab' in which the cysteine residue(s) of the constant domains bear a free thiol group. Antibody fragments originally were produced as pairs of Fab' fragments which have hinge cysteines between them. Other chemical couplings of antibody fragments are also known.

An isolated immunogenically specific paratope or fragment of the antibody is also provided. A specific immunogenic epitope of the antibody can be isolated from the whole antibody by chemical or mechanical disruption of the molecule. The purified fragments thus obtained are tested to determine their immunogenicity and specificity by the methods taught

herein. Immunoreactive paratopes of the antibody, optionally, are synthesized directly. An immunoreactive fragment is defined as an amino acid sequence of at least about two to five consecutive amino acids derived from the antibody amino acid sequence.

One method of producing proteins comprising the antibodies of the present invention is to
5 link two or more peptides or polypeptides together by protein chemistry techniques. For example, peptides or polypeptides can be chemically synthesized using currently available laboratory equipment using either Fmoc (9-fluorenylmethyloxycarbonyl) or Boc (tert -butyloxycarbonyl) chemistry. (Applied Biosystems, Inc., Foster City, CA). One skilled in the art can readily appreciate that a peptide or polypeptide corresponding to the antibody of the present invention, for
10 example, can be synthesized by standard chemical reactions. For example, a peptide or polypeptide can be synthesized and not cleaved from its synthesis resin whereas the other fragment of an antibody can be synthesized and subsequently cleaved from the resin, thereby exposing a terminal group which is functionally blocked on the other fragment. By peptide condensation reactions, these two fragments can be covalently joined via a peptide bond at their carboxyl and
15 amino termini, respectively, to form an antibody, or fragment thereof. (Grant GA (1992) *Synthetic Peptides: A User Guide*. W.H. Freeman and Co., N.Y. (1992); Bodansky M and Trost B., Ed. (1993) *Principles of Peptide Synthesis*. Springer-Verlag Inc., NY. Alternatively, the peptide or polypeptide is independently synthesized in vivo as described above. Once isolated, these independent peptides or polypeptides may be linked to form an antibody or fragment thereof via
20 similar peptide condensation reactions.

For example, enzymatic ligation of cloned or synthetic peptide segments allow relatively short peptide fragments to be joined to produce larger peptide fragments, polypeptides or whole protein domains (Abrahmsen L et al., *Biochemistry*, 30:4151 (1991)). Alternatively, native chemical ligation of synthetic peptides can be utilized to synthetically construct large peptides or
25 polypeptides from shorter peptide fragments. This method consists of a two step chemical reaction (Dawson et al. *Synthesis of Proteins by Native Chemical Ligation*. *Science*, 266:776-779 (1994)). The first step is the chemoselective reaction of an unprotected synthetic peptide-alpha-thioester with another unprotected peptide segment containing an amino-terminal Cys residue to give a thioester-linked intermediate as the initial covalent product. Without a change in the reaction
30 conditions, this intermediate undergoes spontaneous, rapid intramolecular reaction to form a native peptide bond at the ligation site. Application of this native chemical ligation method to the total synthesis of a protein molecule is illustrated by the preparation of human interleukin 8 (IL-8) (Baggiolini M et al. (1992) *FEBS Lett.* 307:97-101; Clark-Lewis I et al., *J.Biol.Chem.*, 269:16075 (1994); Clark-Lewis I et al., *Biochemistry*, 30:3128 (1991); Rajarathnam K et al., *Biochemistry*
35 33:6623-30 (1994)).

Alternatively, unprotected peptide segments are chemically linked where the bond formed between the peptide segments as a result of the chemical ligation is an unnatural (non-peptide) bond (Schnolzer, M et al. Science, 256:221 (1992)). This technique has been used to synthesize analogs of protein domains as well as large amounts of relatively pure proteins with full biological activity (deLisle Milton RC et al., Techniques in Protein Chemistry IV. Academic Press, New York, pp. 257-267 (1992)).

The invention also provides fragments of antibodies which have bioactivity. The polypeptide fragments of the present invention can be recombinant proteins obtained by cloning nucleic acids encoding the polypeptide in an expression system capable of producing the polypeptide fragments thereof, such as an adenovirus or baculovirus expression system. For example, one can determine the active domain of an antibody from a specific hybridoma that can cause a biological effect associated with the interaction of the antibody with EphA2. For example, amino acids found to not contribute to either the activity or the binding specificity or affinity of the antibody can be deleted without a loss in the respective activity. For example, in various embodiments, amino or carboxy-terminal amino acids are sequentially removed from either the native or the modified non-immunoglobulin molecule or the immunoglobulin molecule and the respective activity assayed in one of many available assays. In another example, a fragment of an antibody comprises a modified antibody wherein at least one amino acid has been substituted for the naturally occurring amino acid at a specific position, and a portion of either amino terminal or carboxy terminal amino acids, or even an internal region of the antibody, has been replaced with a polypeptide fragment or other moiety, such as biotin, which can facilitate in the purification of the modified antibody. For example, a modified antibody can be fused to a maltose binding protein, through either peptide chemistry or cloning the respective nucleic acids encoding the two polypeptide fragments into an expression vector such that the expression of the coding region results in a hybrid polypeptide. The hybrid polypeptide can be affinity purified by passing it over an amylose affinity solid support, and the modified antibody receptor can then be separated from the maltose binding region by cleaving the hybrid polypeptide with the specific protease factor Xa. (See, for example, New England Biolabs Product Catalog, 1996, pg. 164.). Similar purification procedures are available for isolating hybrid proteins from eukaryotic cells as well.

The fragments, whether attached to other sequences or not, include insertions, deletions, substitutions, or other selected modifications of particular regions or specific amino acids residues, provided the activity of the fragment is not significantly altered or impaired compared to the nonmodified antibody or antibody fragment. These modifications can provide for some additional property, such as to remove or add amino acids capable of disulfide bonding, to increase its bio-longevity, to alter its secretory characteristics, etc. In any case, the fragment must possess a bioactive property, such as binding activity, regulation of binding at the binding domain, etc.

Functional or active regions of the antibody may be identified by mutagenesis of a specific region of the protein, followed by expression and testing of the expressed polypeptide. Such methods are readily apparent to a skilled practitioner in the art and can include site-specific mutagenesis of the nucleic acid encoding the antigen. (Zoller MJ et al. Nucl. Acids Res. 10:6487-500 (1982).

5 A variety of immunoassay formats may be used to select antibodies that selectively bind with a particular protein, variant, or fragment. For example, solid-phase ELISA immunoassays are routinely used to select antibodies selectively immunoreactive with a protein, protein variant, or fragment thereof. See Harlow and Lane. Antibodies, A Laboratory Manual. Cold Spring Harbor Publications, New York, (1988), for a description of immunoassay formats and conditions that
10 could be used to determine selective binding. The binding affinity of a monoclonal antibody can, for example, be determined by the Scatchard analysis of Munson et al., Anal. Biochem., 107:220 (1980).

Also provided is an antibody reagent kit comprising containers of the monoclonal antibody or fragment thereof of the invention and one or more reagents for detecting binding of the antibody
15 or fragment thereof to the EphA2 receptor molecule. The reagents can include, for example, fluorescent tags, enzymatic tags, or other tags. The reagents can also include secondary or tertiary antibodies or reagents for enzymatic reactions, wherein the enzymatic reactions produce a product that can be visualized.

(b) Functional Nucleic Acids

20 Functional nucleic acids can also be used to regulate the , for example, the function of the disclosed integrins, integrin receptors, molecules that interact with the disclosed integrin receptors, and molecules in the signaling pathways of the disclosed integrin receptors.

Functional nucleic acids are nucleic acid molecules that have a specific function, such as binding a target molecule or catalyzing a specific reaction. Functional nucleic acid molecules can
25 be divided into the following categories, which are not meant to be limiting. For example, functional nucleic acids include antisense molecules, aptamers, ribozymes, triplex forming molecules, and external guide sequences. The functional nucleic acid molecules can act as effectors, inhibitors, modulators, and stimulators of a specific activity possessed by a target molecule, or the functional nucleic acid molecules can possess a de novo activity independent of
30 any other molecules.

Functional nucleic acid molecules can interact with any macromolecule, such as DNA, RNA, polypeptides, or carbohydrate chains. Thus, functional nucleic acids can interact with the mRNA of beta4 integrin for example, or the genomic DNA of alpha6 integrin for example, or they can interact with the polypeptide laminin5, or the gamma2 subunit of laminin5. Often functional
35 nucleic acids are designed to interact with other nucleic acids based on sequence homology

between the target molecule and the functional nucleic acid molecule. In other situations, the specific recognition between the functional nucleic acid molecule and the target molecule is not based on sequence homology between the functional nucleic acid molecule and the target molecule, but rather is based on the formation of tertiary structure that allows specific recognition to take place. Antisense molecules are designed to interact with a target nucleic acid molecule through either canonical or non-canonical base pairing. The interaction of the antisense molecule and the target molecule is designed to promote the destruction of the target molecule through, for example, RNaseH mediated RNA-DNA hybrid degradation. Alternatively the antisense molecule is designed to interrupt a processing function that normally would take place on the target molecule, such as transcription or replication. Antisense molecules can be designed based on the sequence of the target molecule. Numerous methods for optimization of antisense efficiency by finding the most accessible regions of the target molecule exist. Exemplary methods would be in vitro selection experiments and DNA modification studies using DMS and DEPC. It is preferred that antisense molecules bind the target molecule with a dissociation constant (k_d) less than 10^{-6} . It is more preferred that antisense molecules bind with a k_d less than 10^{-8} . It is also more preferred that the antisense molecules bind the target molecule with a k_d less than 10^{-10} . It is also preferred that the antisense molecules bind the target molecule with a k_d less than 10^{-12} . A representative sample of methods and techniques which aid in the design and use of antisense molecules can be found in the following non-limiting list of United States patents: 5,135,917, 5,294,533, 5,627,158, 5,641,754, 5,691,317, 5,780,607, 5,786,138, 5,849,903, 5,856,103, 5,919,772, 5,955,590, 5,990,088, 5,994,320, 5,998,602, 6,005,095, 6,007,995, 6,013,522, 6,017,898, 6,018,042, 6,025,198, 6,033,910, 6,040,296, 6,046,004, 6,046,319, and 6,057,437.

Aptamers are molecules that interact with a target molecule, preferably in a specific way. Typically aptamers are small nucleic acids ranging from 15-50 bases in length that fold into defined secondary and tertiary structures, such as stem-loops or G-quartets. Aptamers can bind small molecules, such as ATP (United States patent 5,631,146) and theophiline (United States patent 5,580,737), as well as large molecules, such as reverse transcriptase (United States patent 5,786,462) and thrombin (United States patent 5,543,293). Aptamers can bind very tightly with k_d s from the target molecule of less than 10^{-12} M. It is preferred that the aptamers bind the target molecule with a k_d less than 10^{-6} . It is more preferred that the aptamers bind the target molecule with a k_d less than 10^{-8} . It is also more preferred that the aptamers bind the target molecule with a k_d less than 10^{-10} . It is also preferred that the aptamers bind the target molecule with a k_d less than 10^{-12} . Aptamers can bind the target molecule with a very high degree of specificity. For example, aptamers have been isolated that have greater than a 10000 fold difference in binding affinities between the target molecule and another molecule that differ at only a single position on the molecule (United States patent 5,543,293). It is preferred that the aptamer have a k_d with the target

molecule at least 10 fold lower than the k_d with a background binding molecule. It is more preferred that the aptamer have a k_d with the target molecule at least 100 fold lower than the k_d with a background binding molecule. It is more preferred that the aptamer have a k_d with the target molecule at least 1000 fold lower than the k_d with a background binding molecule. It is preferred that the aptamer have a k_d with the target molecule at least 10000 fold lower than the k_d with a background binding molecule. It is preferred when doing the comparison for a polypeptide for example, that the background molecule be a different polypeptide. For example, when determining the specificity of beta4 integrin aptamers, the background protein could be bovine serum albumin. Representative examples of how to make and use aptamers to bind a variety of different target molecules can be found in the following non-limiting list of United States patents: 5,476,766, 5,503,978, 5,631,146, 5,731,424, 5,780,228, 5,792,613, 5,795,721, 5,846,713, 5,858,660, 5,861,254, 5,864,026, 5,869,641, 5,958,691, 6,001,988, 6,011,020, 6,013,443, 6,020,130, 6,028,186, 6,030,776, and 6,051,698.

Ribozymes are nucleic acid molecules that are capable of catalyzing a chemical reaction, either intramolecularly or intermolecularly. Ribozymes are thus catalytic nucleic acid. It is preferred that the ribozymes catalyze intermolecular reactions. There are a number of different types of ribozymes that catalyze nuclease or nucleic acid polymerase type reactions which are based on ribozymes found in natural systems, such as hammerhead ribozymes, (for example, but not limited to the following United States patents: 5,334,711, 5,436,330, 5,616,466, 5,633,133, 5,646,020, 5,652,094, 5,712,384, 5,770,715, 5,856,463, 5,861,288, 5,891,683, 5,891,684, 5,985,621, 5,989,908, 5,998,193, 5,998,203, WO 9858058 by Ludwig and Sproat, WO 9858057 by Ludwig and Sproat, and WO 9718312 by Ludwig and Sproat) hairpin ribozymes (for example, but not limited to the following United States patents: 5,631,115, 5,646,031, 5,683,902, 5,712,384, 5,856,188, 5,866,701, 5,869,339, and 6,022,962), and tetrahymena ribozymes (for example, but not limited to the following United States patents: 5,595,873 and 5,652,107). There are also a number of ribozymes that are not found in natural systems, but which have been engineered to catalyze specific reactions de novo (for example, but not limited to the following United States patents: 5,580,967, 5,688,670, 5,807,718, and 5,910,408). Preferred ribozymes cleave RNA or DNA substrates, and more preferably cleave RNA substrates. Ribozymes typically cleave nucleic acid substrates through recognition and binding of the target substrate with subsequent cleavage. This recognition is often based mostly on canonical or non-canonical base pair interactions. This property makes ribozymes particularly good candidates for target specific cleavage of nucleic acids because recognition of the target substrate is based on the target substrates sequence. Representative examples of how to make and use ribozymes to catalyze a variety of different reactions can be found in the following non-limiting list of United States patents: 5,646,042, 5,693,535, 5,731,295, 5,811,300, 5,837,855, 5,869,253, 5,877,021, 5,877,022, 5,972,699,

5,972,704, 5,989,906, and 6,017,756.

Triplex forming functional nucleic acid molecules are molecules that can interact with either double-stranded or single-stranded nucleic acid. When triplex molecules interact with a target region, a structure called a triplex is formed, in which there are three strands of DNA forming a complex dependant on both Watson-Crick and Hoogsteen base-pairing. Triplex molecules are preferred because they can bind target regions with high affinity and specificity. It is preferred that the triplex forming molecules bind the target molecule with a k_d less than 10^{-6} . It is more preferred that the triplex forming molecules bind with a k_d less than 10^{-8} . It is also more preferred that the triplex forming molecules bind the target molecule with a k_d less than 10^{-10} . It is also preferred that the triplex forming molecules bind the target molecule with a k_d less than 10^{-12} . Representative examples of how to make and use triplex forming molecules to bind a variety of different target molecules can be found in the following non-limiting list of United States patents: 5,176,996, 5,645,985, 5,650,316, 5,683,874, 5,693,773, 5,834,185, 5,869,246, 5,874,566, and 5,962,426.

External guide sequences (EGSs) are molecules that bind a target nucleic acid molecule forming a complex, and this complex is recognized by RNase P, which cleaves the target molecule. EGSs can be designed to specifically target a RNA molecule of choice. RNase P aids in processing transfer RNA (tRNA) within a cell. Bacterial RNase P can be recruited to cleave virtually any RNA sequence by using an EGS that causes the target RNA:EGS complex to mimic the natural tRNA substrate. (WO 92/03566 by Yale, and Forster and Altman, Science 238:407-409 (1990)).

Similarly, eukaryotic EGS/RNase P-directed cleavage of RNA can be utilized to cleave desired targets within eukaryotic cells. (Yuan et al., Proc. Natl. Acad. Sci. USA 89:8006-8010 (1992); WO 93/22434 by Yale; WO 95/24489 by Yale; Yuan and Altman, EMBO J 14:159-168 (1995), and Carrara et al., Proc. Natl. Acad. Sci. (USA) 92:2627-2631 (1995)). Representative examples of how to make and use EGS molecules to facilitate cleavage of a variety of different target molecules be found in the following non-limiting list of United States patents: 5,168,053, 5,624,824, 5,683,873, 5,728,521, 5,869,248, and 5,877,162

(c) Small molecules

Small molecules can also be used to regulate, for example, the function of the disclosed integrins, integrin receptors, molecules that interact with the disclosed integrin receptors, and molecules in the signaling pathways of the disclosed integrin receptors. Those of skill in the art understand how to generate small molecules of this type, and exemplary libraries and methods for isolating small molecule regulators are disclosed herein.

b) Compositions identified by screening with disclosed compositions / combinatorial chemistry

(1) Combinatorial chemistry

The disclosed compositions can be used as targets for any combinatorial technique to
5 identify molecules or macromolecular molecules that interact with the disclosed compositions,
such as beta4 integrin, alpha6 integrin, or the gamma2 subunit of laminin5, in a desired way. The
nucleic acids, peptides, and related molecules disclosed herein can be used as targets for the
combinatorial approaches. Also disclosed are the compositions that are identified through
combinatorial techniques or screening techniques in which the herein disclosed compositions, for
10 example set forth in SEQ ID NOS:1-19 or portions thereof, are used as the target or reagent in a
combinatorial or screening protocol.

It is understood that when using the disclosed compositions in combinatorial techniques or
screening methods, molecules, such as macromolecular molecules, will be identified that have
particular desired properties such as inhibition or stimulation or the target molecule's function.
15 The molecules identified and isolated when using the disclosed compositions, such as, beta4
integrin, alpha6 integrin, or the gamma2 subunit of laminin5, are also disclosed. Thus, the products
produced using the combinatorial or screening approaches that involve the disclosed compositions,
such as, beta4 integrin, alpha6 integrin, or the gamma2 subunit of laminin5, are also considered
herein disclosed.

20 Combinatorial chemistry includes but is not limited to all methods for isolating small
molecules or macromolecules that are capable of binding either a small molecule or another
macromolecule, typically in an iterative process. Proteins, oligonucleotides, and sugars are
examples of macromolecules. For example, oligonucleotide molecules with a given function,
catalytic or ligand-binding, can be isolated from a complex mixture of random oligonucleotides in
25 what has been referred to as "in vitro genetics" (Szostak, TIBS 19:89, 1992). One synthesizes a
large pool of molecules bearing random and defined sequences and subjects that complex mixture,
for example, approximately 10^{15} individual sequences in 100 μ g of a 100 nucleotide RNA, to some
selection and enrichment process. Through repeated cycles of affinity chromatography and PCR
amplification of the molecules bound to the ligand on the solid support, Ellington and Szostak
30 (1990) estimated that 1 in 10^{10} RNA molecules folded in such a way as to bind a small molecule
dyes. DNA molecules with such ligand-binding behavior have been isolated as well (Ellington and
Szostak, 1992; Bock et al, 1992). Techniques aimed at similar goals exist for small organic
molecules, proteins, antibodies and other macromolecules known to those of skill in the art.
Screening sets of molecules for a desired activity whether based on small organic libraries,
35 oligonucleotides, or antibodies is broadly referred to as combinatorial chemistry. Combinatorial
techniques are particularly suited for defining binding interactions between molecules and for

isolating molecules that have a specific binding activity, often called aptamers when the macromolecules are nucleic acids.

There are a number of methods for isolating proteins which either have de novo activity or a modified activity. For example, phage display libraries have been used to isolate numerous
5 peptides that interact with a specific target. (See for example, United States Patent No. 6,031,071; 5,824,520; 5,596,079; and 5,565,332 which are herein incorporated by reference at least for their material related to phage display and methods relate to combinatorial chemistry)

A preferred method for isolating proteins that have a given function is described by Roberts and Szostak (Roberts R.W. and Szostak J.W. Proc. Natl. Acad. Sci. USA,
10 94(23)12997-302 (1997). This combinatorial chemistry method couples the functional power of proteins and the genetic power of nucleic acids. An RNA molecule is generated in which a puromycin molecule is covalently attached to the 3'-end of the RNA molecule. An *in vitro* translation of this modified RNA molecule causes the correct protein, encoded by the RNA to be translated. In addition, because of the attachment of the puromycin, a peptidyl acceptor which
15 cannot be extended, the growing peptide chain is attached to the puromycin which is attached to the RNA. Thus, the protein molecule is attached to the genetic material that encodes it. Normal *in vitro* selection procedures can now be done to isolate functional peptides. Once the selection procedure for peptide function is complete traditional nucleic acid manipulation procedures are performed to amplify the nucleic acid that codes for the selected functional peptides. After
20 amplification of the genetic material, new RNA is transcribed with puromycin at the 3'-end, new peptide is translated and another functional round of selection is performed. Thus, protein selection can be performed in an iterative manner just like nucleic acid selection techniques. The peptide which is translated is controlled by the sequence of the RNA attached to the puromycin. This sequence can be anything from a random sequence engineered for optimum translation (i.e.
25 no stop codons etc.) or it can be a degenerate sequence of a known RNA molecule to look for improved or altered function of a known peptide. The conditions for nucleic acid amplification and *in vitro* translation are well known to those of ordinary skill in the art and are preferably performed as in Roberts and Szostak (Roberts R.W. and Szostak J.W. Proc. Natl. Acad. Sci. USA, 94(23)12997-302 (1997)).

30 Another preferred method for combinatorial methods designed to isolate peptides is described in Cohen et al. (Cohen B.A., et al., Proc. Natl. Acad. Sci. USA 95(24):14272-7 (1998)). This method utilizes and modifies two-hybrid technology. Yeast two-hybrid systems are useful for the detection and analysis of protein:protein interactions. The two-hybrid system, initially described in the yeast *Saccharomyces cerevisiae*, is a powerful molecular genetic
35 technique for identifying new regulatory molecules, specific to the protein of interest (Fields and

Song, *Nature* 340:245-6 (1989)). Cohen et al., modified this technology so that novel interactions between synthetic or engineered peptide sequences could be identified which bind a molecule of choice. The benefit of this type of technology is that the selection is done in an intracellular environment. The method utilizes a library of peptide molecules that attached to an acidic
 5 activation domain. A peptide of choice, for example a portion of beta4 or alpha6 or gamma2 or laminin5 is attached to a DNA binding domain of a transcriptional activation protein, such as Gal 4. By performing the Two-hybrid technique on this type of system, molecules that bind the portion of beta4 or alpha6 or gamma2 or laminin5 can be identified.

There are molecules that can act like antibodies, in that they can having varying binding
 10 specificities, that are based on a fibronectin motif. The fibronectin type III domain (FN3) is a small autonomous folding unit. This FN3 domain can be found in numerous proteins that bind ligand, such as animal proteins. The beta-sandwich structure of FN3 closely resembles that of immunoglobulin domains. FN3 mutants can be isolated using combinatorial approaches disclosed herein, for example phage display, that bind desired targets. Typically the libraries of FN3
 15 molecules have been randomized in the two surface loops. Thus, FN3 can be used at least as a scaffold for engineering novel binding proteins. (Koide A, Bailey CW, Huang X, Koide S., "The fibronectin type III domain as a scaffold for novel binding proteins." *J Mol Biol* 1998: 284,1141-1151 which is herein incorporated by reference at least for material related to the fibronectin based novel binding proteins).

20 Using methodology well known to those of skill in the art, in combination with various combinatorial libraries, one can isolate and characterize those small molecules or macromolecules, which bind to or interact with the desired target. The relative binding affinity of these compounds can be compared and optimum compounds identified using competitive binding studies, which are well known to those of skill in the art.

25 Techniques for making combinatorial libraries and screening combinatorial libraries to isolate molecules which bind a desired target are well known to those of skill in the art. Representative techniques and methods can be found in but are not limited to United States patents 5,084,824, 5,288,514, 5,449,754, 5,506,337, 5,539,083, 5,545,568, 5,556,762, 5,565,324, 5,565,332, 5,573,905, 5,618,825, 5,619,680, 5,627,210, 5,646,285, 5,663,046, 5,670,326,
 30 5,677,195, 5,683,899, 5,688,696, 5,688,997, 5,698,685, 5,712,146, 5,721,099, 5,723,598, 5,741,713, 5,792,431, 5,807,683, 5,807,754, 5,821,130, 5,831,014, 5,834,195, 5,834,318, 5,834,588, 5,840,500, 5,847,150, 5,856,107, 5,856,496, 5,859,190, 5,864,010, 5,874,443, 5,877,214, 5,880,972, 5,886,126, 5,886,127, 5,891,737, 5,916,899, 5,919,955, 5,925,527, 5,939,268, 5,942,387, 5,945,070, 5,948,696, 5,958,702, 5,958,792, 5,962,337, 5,965,719,
 35 5,972,719, 5,976,894, 5,980,704, 5,985,356, 5,999,086, 6,001,579, 6,004,617, 6,008,321,

6,017,768, 6,025,371, 6,030,917, 6,040,193, 6,045,671, 6,045,755, 6,060,596, and 6,061,636.

Combinatorial libraries can be made from a wide array of molecules using a number of different synthetic techniques. For example, libraries containing fused 2,4-pyrimidinediones (United States patent 6,025,371) dihydrobenzopyrans (United States Patent 6,017,768 and
5 5,821,130), amide alcohols (United States Patent 5,976,894), hydroxy-amino acid amides (United States Patent 5,972,719) carbohydrates (United States patent 5,965,719), 1,4-benzodiazepin-2,5-diones (United States patent 5,962,337), cyclics (United States patent 5,958,792), biaryl amino acid amides (United States patent 5,948,696), thiophenes (United States patent 5,942,387), tricyclic Tetrahydroquinolines (United States patent 5,925,527), benzofurans (United States patent
10 5,919,955), isoquinolines (United States patent 5,916,899), hydantoin and thiohydantoin (United States patent 5,859,190), indoles (United States patent 5,856,496), imidazol-pyrido-indole and imidazol-pyrido-benzothiophenes (United States patent 5,856,107) substituted 2-methylene-2, 3-dihydrothiazoles (United States patent 5,847,150), quinolines (United States patent 5,840,500), PNA (United States patent 5,831,014), containing tags (United States patent 5,721,099),
15 polyketides (United States patent 5,712,146), morpholino-subunits (United States patent 5,698,685 and 5,506,337), sulfamides (United States patent 5,618,825), and benzodiazepines (United States patent 5,288,514).

Screening molecules similar to alpha6 for inhibition of alpha6beta4 formation is a method of identifying and isolating desired compounds that can inhibit the formation of A6B4 receptor.
20 For example, the disclosed compositions, such as alpha6 integrin or beta4 integrin can be used as targets in a selection scheme disclosed herein, and then the counter part integrin could be used as a competitive inhibitor to isolate the desired molecules. For example, a library of molecules could be incubated with beta4 integrin, which is bound to a solid support. The solid support can be washed to remove the unbound molecules and then the solid support can be incubated with, for
25 example, alpha6 integrin at a concentration that will saturate all beta4 binding sites. The molecules which are collected in the flowthrough after washing the solid support will be enriched for molecules that interact with beta4 integrin in a way that is competitive to the alpha6-beta4 interaction. Likewise, the solid support, bound with a target integrin, or more preferably a target integrin receptor, such as alpha6beta4 receptor, could also be washed, with for example, laminin5
30 or the gamma2 subunit of laminin5 at a concentration that will saturate all of the gamma2 binding sites on the beta4 integrin. Collection of the wash under these conditions will yield a population of molecules enriched for molecules that competitively interact with beta4 integrin at the beta4-gamma2 site. Another example, is the following: bind target to solid support on microtiter plate. Incubate with ligand in presence of gridded subset of library members (or single compounds),
35 wash, identify competitor by reduction of ligand binding. It is understood that the exemplary discussions of alpha6beta4 and/or beta4 are equally applicable to alpha6, as well as other

alpha6betax receptors, such alpha6beta1.

Also disclosed are methods of isolating molecules that bind with a target molecule selected from the group consisting, B4 integrin, alpha6 integrin, and the gamma2 subunit of laminin5 comprising 1) contacting a library of molecules with the target molecule and 2)
5 collecting molecules that bind the target molecule producing an enriched population of molecules.

Disclosed are methods, further comprising the step of repeating steps 1 and 2 with the enriched population of molecules, and/or wherein the library comprises a small molecule, peptide, peptide mimetic, or oligonucleotide.

As used herein combinatorial methods and libraries included traditional screening methods
10 and libraries as well as methods and libraries used in iterative processes.

(2) Computer assisted design

The disclosed compositions can be used as targets for any molecular modeling technique to identify either the structure of the disclosed compositions or to identify potential or actual molecules, such as small molecules, which interact in a desired way with the disclosed
15 compositions. The nucleic acids, peptides, and related molecules disclosed herein can be used as targets in any molecular modeling program or approach.

It is understood that when using the disclosed compositions in modeling techniques, molecules, such as macromolecular molecules, will be identified that have particular desired properties such as inhibition or stimulation or the target molecule's function. The molecules
20 identified and isolated when using the disclosed compositions, such as, beta4, are also disclosed. Thus, the products produced using the molecular modeling approaches that involve the disclosed compositions, such as, beta4 integrin, alpha6 integrin, or laminin5, are also considered herein disclosed.

Thus, one way to isolate molecules that bind a molecule of choice is through rational
25 design. This is achieved through structural information and computer modeling. Computer modeling technology allows visualization of the three-dimensional atomic structure of a selected molecule and the rational design of new compounds that will interact with the molecule. The three-dimensional construct typically depends on data from x-ray crystallographic analyses or NMR imaging of the selected molecule. The molecular dynamics require force field data. The
30 computer graphics systems enable prediction of how a new compound will link to the target molecule and allow experimental manipulation of the structures of the compound and target molecule to perfect binding specificity. Prediction of what the molecule-compound interaction will be when small changes are made in one or both requires molecular mechanics software and computationally intensive computers, usually coupled with user-friendly, menu-driven interfaces

between the molecular design program and the user.

Examples of molecular modeling systems are the CHARMM and QUANTA programs, Polygen Corporation, Waltham, MA. CHARMM performs the energy minimization and molecular dynamics functions. QUANTA performs the construction, graphic modeling and analysis of
5 molecular structure. QUANTA allows interactive construction, modification, visualization, and analysis of the behavior of molecules with each other.

A number of articles review computer modeling of drugs interactive with specific proteins, such as Rotivinen, et al., 1988 *Acta Pharmaceutica Fennica* 97, 159-166; Ripka, *New Scientist* 54-57 (June 16, 1988); McKinaly and Rossmann, 1989 *Annu. Rev. Pharmacol. Toxicol.* 29, 111-
10 122; Perry and Davies, QSAR: Quantitative Structure-Activity Relationships in Drug Design pp. 189-193 (Alan R. Liss, Inc. 1989); Lewis and Dean, 1989 *Proc. R. Soc. Lond.* 236, 125-140 and 141-162; and, with respect to a model enzyme for nucleic acid components, Askew, et al., 1989 *J. Am. Chem. Soc.* 111, 1082-1090. Other computer programs that screen and graphically depict chemicals are available from companies such as BioDesign, Inc., Pasadena, CA., Allelix,
15 Inc, Mississauga, Ontario, Canada, and Hypercube, Inc., Cambridge, Ontario. Although these are primarily designed for application to drugs specific to particular proteins, they can be adapted to design of molecules specifically interacting with specific regions of DNA or RNA, once that region is identified.

Although described above with reference to design and generation of compounds which
20 could alter binding, one could also screen libraries of known compounds, including natural products or synthetic chemicals, and biologically active materials, including proteins, for compounds which alter substrate binding or enzymatic activity.

c) Nucleic acids

There are a variety of molecules disclosed herein that are nucleic acid based, including for
25 example the nucleic acids that encode, for example beta4 and the gamma2 subunit of laminin5, as well as various functional nucleic acids. The disclosed nucleic acids are made up of for example, nucleotides, nucleotide analogs, or nucleotide substitutes. Non-limiting examples of these and other molecules are discussed herein. It is understood that for example, when a vector is expressed in a cell, that the expressed mRNA will typically be made up of A, C, G, and U. Likewise, it is
30 understood that if, for example, an antisense molecule is introduced into a cell or cell environment through for example exogenous delivery, it is advantageous that the antisense molecule be made up of nucleotide analogs that reduce the degradation of the antisense molecule in the cellular environment.

(1) Nucleotides and related molecules

35 A nucleotide is a molecule that contains a base moiety, a sugar moiety and a phosphate

moiety. Nucleotides can be linked together through their phosphate moieties and sugar moieties creating an internucleoside linkage. The base moiety of a nucleotide can be adenin-9-yl (A), cytosin-1-yl (C), guanin-9-yl (G), uracil-1-yl (U), and thymine-1-yl (T). The sugar moiety of a nucleotide is a ribose or a deoxyribose. The phosphate moiety of a nucleotide is pentavalent
5 phosphate. An non-limiting example of a nucleotide would be 3'-AMP (3'-adenosine monophosphate) or 5'-GMP (5'-guanosine monophosphate).

A nucleotide analog is a nucleotide which contains some type of modification to either the base, sugar, or phosphate moieties. Modifications to the base moiety would include natural and synthetic modifications of A, C, G, and T/U as well as different purine or pyrimidine bases, such
10 as uracil-5-yl (.psi.), hypoxanthin-9-yl (I), and 2-aminoadenin-9-yl. A modified base includes but is not limited to 5-methylcytosine (5-me-C), 5-hydroxymethyl cytosine, xanthine, hypoxanthine, 2-aminoadenine, 6-methyl and other alkyl derivatives of adenine and guanine, 2-propyl and other alkyl derivatives of adenine and guanine, 2-thiouracil, 2-thiothymine and 2-thiocytosine, 5-halouracil and cytosine, 5-propynyl uracil and cytosine, 6-azo uracil, cytosine and thymine, 5-
15 uracil (pseudouracil), 4-thiouracil, 8-halo, 8-amino, 8-thiol, 8-thioalkyl, 8-hydroxyl and other 8-substituted adenines and guanines, 5-halo particularly 5-bromo, 5-trifluoromethyl and other 5-substituted uracils and cytosines, 7-methylguanine and 7-methyladenine, 8-azaguanine and 8-azaadenine, 7-deazaguanine and 7-deazaadenine and 3-deazaguanine and 3-deazaadenine. Additional base modifications can be found for example in U.S. Pat. No. 3,687,808, Englisch et
20 al., *Angewandte Chemie, International Edition*, 1991, 30, 613, and Sanghvi, Y. S., Chapter 15, *Antisense Research and Applications*, pages 289-302, Crooke, S. T. and Lebleu, B. ed., CRC Press, 1993. Certain nucleotide analogs, such as 5-substituted pyrimidines, 6-azapyrimidines and N-2, N-6 and O-6 substituted purines, including 2-aminopropyladenine, 5-propynyluracil and 5-propynylcytosine. 5-methylcytosine can increase the stability of duplex formation. Base
25 modifications can be combined, for example, with a sugar modification, such as 2'-O-methoxyethyl, to achieve unique properties such as increased duplex stability. There are numerous United States patents such as 4,845,205; 5,130,302; 5,134,066; 5,175,273; 5,367,066; 5,432,272; 5,457,187; 5,459,255; 5,484,908; 5,502,177; 5,525,711; 5,552,540; 5,587,469; 5,594,121, 5,596,091; 5,614,617; and 5,681,941, which detail and describe a range of base
30 modifications. Each of these patents is herein incorporated by reference.

Nucleotide analogs can also include modifications of the sugar moiety. Modifications to the sugar moiety would include natural modifications of the ribose and deoxy ribose as well as synthetic modifications. Sugar modifications include but are not limited to the following modifications at the 2' position: OH-; F-; O-, S-, or N-alkyl; O-, S-, or N-alkenyl; O-, S- or N-
35 alkynyl; or O-alkyl-O-alkyl, wherein the alkyl, alkenyl and alkynyl may be substituted or unsubstituted C₁ to C₁₀, alkyl or C₂ to C₁₀ alkenyl and alkynyl. 2' sugar modifications also include

but are not limited to $-O[(CH_2)_n O]_m CH_3$, $-O(CH_2)_n OCH_3$, $-O(CH_2)_n NH_2$, $-O(CH_2)_n CH_3$, $-O(CH_2)_n -ONH_2$, and $-O(CH_2)_n ON[(CH_2)_n CH_3]_2$, where n and m are from 1 to about 10.

Other modifications at the 2' position include but are not limited to: C_1 to C_{10} lower alkyl, substituted lower alkyl, alkaryl, aralkyl, O-alkaryl or O-aralkyl, SH, SCH_3 , OCN, Cl, Br, CN, CF_3 , OCF_3 , $SOCH_3$, $SO_2 CH_3$, ONO_2 , NO_2 , N_3 , NH_2 , heterocycloalkyl, heterocycloalkaryl, aminoalkylamino, polyalkylamino, substituted silyl, an RNA cleaving group, a reporter group, an intercalator, a group for improving the pharmacokinetic properties of an oligonucleotide, or a group for improving the pharmacodynamic properties of an oligonucleotide, and other substituents having similar properties. Similar modifications may also be made at other positions on the sugar, particularly the 3' position of the sugar on the 3' terminal nucleotide or in 2'-5' linked oligonucleotides and the 5' position of 5' terminal nucleotide. Modified sugars would also include those that contain modifications at the bridging ring oxygen, such as CH_2 and S. Nucleotide sugar analogs may also have sugar mimetics such as cyclobutyl moieties in place of the pentofuranosyl sugar. There are numerous United States patents that teach the preparation of such modified sugar structures such as 4,981,957; 5,118,800; 5,319,080; 5,359,044; 5,393,878; 5,446,137; 5,466,786; 5,514,785; 5,519,134; 5,567,811; 5,576,427; 5,591,722; 5,597,909; 5,610,300; 5,627,053; 5,639,873; 5,646,265; 5,658,873; 5,670,633; and 5,700,920, each of which is herein incorporated by reference in its entirety.

Nucleotide analogs can also be modified at the phosphate moiety. Modified phosphate moieties include but are not limited to those that can be modified so that the linkage between two nucleotides contains a phosphorothioate, chiral phosphorothioate, phosphorodithioate, phosphotriester, aminoalkylphosphotriester, methyl and other alkyl phosphonates including 3'-alkylene phosphonate and chiral phosphonates, phosphinates, phosphoramidates including 3'-amino phosphoramidate and aminoalkylphosphoramidates, thionophosphoramidates, thionoalkylphosphonates, thionoalkylphosphotriesters, and boranophosphates. It is understood that these phosphate or modified phosphate linkage between two nucleotides can be through a 3'-5' linkage or a 2'-5' linkage, and the linkage can contain inverted polarity such as 3'-5' to 5'-3' or 2'-5' to 5'-2'. Various salts, mixed salts and free acid forms are also included. Numerous United States patents teach how to make and use nucleotides containing modified phosphates and include but are not limited to, 3,687,808; 4,469,863; 4,476,301; 5,023,243; 5,177,196; 5,188,897; 5,264,423; 5,276,019; 5,278,302; 5,286,717; 5,321,131; 5,399,676; 5,405,939; 5,453,496; 5,455,233; 5,466,677; 5,476,925; 5,519,126; 5,536,821; 5,541,306; 5,550,111; 5,563,253; 5,571,799; 5,587,361; and 5,625,050, each of which is herein incorporated by reference.

It is understood that nucleotide analogs need only contain a single modification, but may also contain multiple modifications within one of the moieties or between different moieties.

Nucleotide substitutes are molecules having similar functional properties to nucleotides, but which do not contain a phosphate moiety, such as peptide nucleic acid (PNA). Nucleotide substitutes are molecules that will recognize nucleic acids in a Watson-Crick or Hoogsteen manner, but which are linked together through a moiety other than a phosphate moiety.

- 5 Nucleotide substitutes are able to conform to a double helix type structure when interacting with the appropriate target nucleic acid.

Nucleotide substitutes are nucleotides or nucleotide analogs that have had the phosphate moiety and/or sugar moieties replaced. Nucleotide substitutes do not contain a standard phosphorus atom. Substitutes for the phosphate can be for example, short chain alkyl or
 10 cycloalkyl internucleoside linkages, mixed heteroatom and alkyl or cycloalkyl internucleoside linkages, or one or more short chain heteroatomic or heterocyclic internucleoside linkages. These include those having morpholino linkages (formed in part from the sugar portion of a nucleoside); siloxane backbones; sulfide, sulfoxide and sulfone backbones; formacetyl and thioformacetyl backbones; methylene formacetyl and thioformacetyl backbones; alkene containing backbones;
 15 sulfamate backbones; methyleneimino and methylenehydrazino backbones; sulfonate and sulfonamide backbones; amide backbones; and others having mixed N, O, S and CH₂ component parts. Numerous United States patents disclose how to make and use these types of phosphate replacements and include but are not limited to 5,034,506; 5,166,315; 5,185,444; 5,214,134; 5,216,141; 5,235,033; 5,264,562; 5,264,564; 5,405,938; 5,434,257; 5,466,677; 5,470,967;
 20 5,489,677; 5,541,307; 5,561,225; 5,596,086; 5,602,240; 5,610,289; 5,602,240; 5,608,046; 5,610,289; 5,618,704; 5,623,070; 5,663,312; 5,633,360; 5,677,437; and 5,677,439, each of which is herein incorporated by reference.

It is also understood in a nucleotide substitute that both the sugar and the phosphate moieties of the nucleotide can be replaced, by for example an amide type linkage
 25 (aminoethylglycine) (PNA). United States patents 5,539,082; 5,714,331; and 5,719,262 teach how to make and use PNA molecules, each of which is herein incorporated by reference. (See also Nielsen et al., Science, 1991, 254, 1497-1500).

It is also possible to link other types of molecules (conjugates) to nucleotides or nucleotide analogs to enhance for example, cellular uptake. Conjugates can be chemically linked to the
 30 nucleotide or nucleotide analogs. Such conjugates include but are not limited to lipid moieties such as a cholesterol moiety (Letsinger et al., Proc. Natl. Acad. Sci. USA, 1989,

86, 6553-6556), cholic acid (Manoharan et al., Bioorg. Med. Chem. Let., 1994, 4, 1053-1060), a thioether, e.g., hexyl-S-tritylthiol (Manoharan et al., Ann. N.Y. Acad. Sci., 1992, 660, 306-309; Manoharan et al., Bioorg. Med. Chem. Let., 1993, 3, 2765-2770), a thiocholesterol
 35 (Oberhauser et al., Nucl. Acids Res., 1992, 20, 533-538), an aliphatic chain, e.g., dodecandiol or

undecyl residues (Saison-Behmoaras et al., EMBO J., 1991, 10, 1111-1118; Kabanov et al., FEBS Lett., 1990, 259, 327-330; Svinarchuk et al., Biochimie, 1993, 75, 49-54), a phospholipid, e.g., dihexadecyl-rac-glycerol or triethylammonium 1,2-di-O-hexadecyl-rac-glycero-3-H-phosphonate (Manoharan et al., Tetrahedron Lett., 1995, 36, 3651-3654; Shea et al., Nucl. Acids Res., 1990, 18, 3777-3783), a polyamine or a polyethylene glycol chain (Manoharan et al., Nucleosides & Nucleotides, 1995, 14, 969-973), or adamantane acetic acid (Manoharan et al., Tetrahedron Lett., 1995, 36, 3651-3654), a palmityl moiety (Mishra et al., Biochim. Biophys. Acta, 1995, 1264, 229-237), or an octadecylamine or hexylamino-carbonyl-oxysterol moiety (Crooke et al., J. Pharmacol. Exp. Ther., 1996, 277, 923-937. Numerous United States patents teach the preparation of such conjugates and include, but are not limited to U.S. Pat. Nos. 4,828,979; 4,948,882; 5,218,105; 5,525,465; 5,541,313; 5,545,730; 5,552,538; 5,578,717; 5,580,731; 5,580,731; 5,591,584; 5,109,124; 5,118,802; 5,138,045; 5,414,077; 5,486,603; 5,512,439; 5,578,718; 5,608,046; 4,587,044; 4,605,735; 4,667,025; 4,762,779; 4,789,737; 4,824,941; 4,835,263; 4,876,335; 4,904,582; 4,958,013; 5,082,830; 5,112,963; 5,214,136; 5,082,830; 5,112,963; 5,214,136; 5,245,022; 5,254,469; 5,258,506; 5,262,536; 5,272,250; 5,292,873; 5,317,098; 5,371,241; 5,391,723; 5,416,203; 5,451,463; 5,510,475; 5,512,667; 5,514,785; 5,565,552; 5,567,810; 5,574,142; 5,585,481; 5,587,371; 5,595,726; 5,597,696; 5,599,923; 5,599,928 and 5,688,941, each of which is herein incorporated by reference.

It is understood that an oligonucleotide can be made from any combination of nucleotides, nucleotide analogs, or nucleotide substitutes disclosed herein or related molecules not specifically recited herein.

A Watson-Crick interaction is at least one interaction with the Watson-Crick face of a nucleotide, nucleotide analog, or nucleotide substitute. The Watson-Crick face of a nucleotide, nucleotide analog, or nucleotide substitute includes the C2, N1, and C6 positions of a purine based nucleotide, nucleotide analog, or nucleotide substitute and the C2, N3, C4 positions of a pyrimidine based nucleotide, nucleotide analog, or nucleotide substitute.

A Hoogsteen interaction is the interaction that takes place on the Hoogsteen face of a nucleotide or nucleotide analog, which is exposed in the major groove of duplex DNA. The Hoogsteen face includes the N7 position and reactive groups (NH₂ or O) at the C6 position of purine nucleotides.

(2) Sequences

There are a variety of sequences related to the beta4 integrin or the laminin5-gamma2 gene or the alpha6 integrin gene having the following Genbank Accession Numbers 6453379, 4557674, and AH006634, respectively. These sequences and others are herein incorporated by reference in their entireties as well as for individual subsequences contained therein.

One particular sequence set forth in SEQ ID NO:1 (beta4 integrin) and having Genbank accession number 6453379 is used herein, at various points, as an example, to exemplify the disclosed compositions and methods (or when another particular sequence is used as an example). It is understood that the description related to this sequence is applicable to any sequence related to
5 beta4 integrin or any of the other molecules disclosed herein, such as alpha6 integrin, or the subunits of laminin5, such as gamma2, unless specifically indicated otherwise. Those of skill in the art understand how to resolve sequence discrepancies and differences and to adjust the compositions and methods relating to a particular sequence to other related sequences (i.e. sequences of beta4 integrin). Primers and/or probes can be designed for any beta4 integrin
10 sequence given the information disclosed herein and known in the art.

(3) Primers and probes

Disclosed are compositions including primers and probes, which are capable of interacting with the for example, the alpha6 gene or mRNA, beta4 gene or mRNA, or gamma2 subunit of the laminin5 ligand gene as disclosed herein or mRNA as wells as primers or probes for any of the
15 sequences or fragments of the sequences, set forth in SEQ ID NOs:1,3,5,7,9,11, and 13. In certain embodiments the primers are used to support DNA amplification reactions. Typically the primers will be capable of being extended in a sequence specific manner. Extension of a primer in a sequence specific manner includes any methods wherein the sequence and/or composition of the nucleic acid molecule to which the primer is hybridized or otherwise associated directs or
20 influences the composition or sequence of the product produced by the extension of the primer. Extension of the primer in a sequence specific manner therefore includes, but is not limited to, PCR, DNA sequencing, DNA extension, DNA polymerization, RNA transcription, or reverse transcription. Techniques and conditions that amplify the primer in a sequence specific manner are preferred. In certain embodiments the primers are used for the DNA amplification reactions,
25 such as PCR or direct sequencing. It is understood that in certain embodiments the primers can also be extended using non-enzymatic techniques, where for example, the nucleotides or oligonucleotides used to extend the primer are modified such that they will chemically react to extend the primer in a sequence specific manner. Typically the disclosed primers hybridize with the beta4 gene, alpha6 gene, or laminin5-gamma2 subunit gene, for example, or region of the
30 beta4 gene, alpha6 gene, or laminin5-gamma2 subunit gene, for example, or they hybridize with the complement of the beta4 gene, alpha6 gene, or laminin5-gamma2 subunit gene, for example, or complement of a region of the beta4 gene, alpha6 gene, or laminin5-gamma2 subunit gene, or any of the sequences or fragments of the sequences, set forth in SEQ ID NOs:1,3,5,7,9,11, and 13, for example.

35 The size of the primers or probes for interaction with the beta4 gene, alpha6 gene, or laminin5-gamma2 subunit gene, for example, in certain embodiments can be any size that supports

the desired enzymatic manipulation of the primer, such as DNA amplification or the simple hybridization of the probe or primer. A typical primer or probe for beta4 gene, alpha6 gene, or laminin5-gamma2 subunit gene, for example, or primer or probe for any of the sequences or fragments of the sequences, set forth in SEQ ID NOs:1,3,5,7,9,11, and 13 would be at least about

5 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, 20, 21, 22, 23, 24, 25, 26, 27, 28, 29, 30, 31, 32, 33, 34, 35, 36, 37, 38, 39, 40, 41, 42, 43, 44, 45, 46, 47, 48, 49, 50, 51, 52, 53, 54, 55, 56, 57, 58, 59, 60, 61, 62, 63, 64, 65, 66, 67, 68, 69, 70, 71, 72, 73, 74, 75, 76, 77, 78, 79, 80, 81, 82, 83, 84, 85, 86, 87, 88, 89, 90, 91, 92, 93, 94, 95, 96, 97, 98, 99, 100, 125, 150, 175, 200, 225, 250, 275, 300, 325, 350, 375, 400, 425, 450, 475, 500, 550, 600, 650, 700, 750, 800, 850, 900, 950, 1000, 1250, 1500, 1750, 2000, 2250, 2500, 2750, 3000, 3500, or 4000 nucleotides long.

In other embodiments a primer or probe for beta4 gene, alpha6 gene, or laminin5-gamma2 subunit gene, for example, primer or probe or a primer or probe for any of the sequences or fragments of the sequences, set forth in SEQ ID NOs:1,3,5,7,9,11, and 13 can be less than or equal to about 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, 20, 21, 22, 23, 24, 25, 26, 27, 28, 29, 30,

15 31, 32, 33, 34, 35, 36, 37, 38, 39, 40, 41, 42, 43, 44, 45, 46, 47, 48, 49, 50, 51, 52, 53, 54, 55, 56, 57, 58, 59, 60, 61, 62, 63, 64, 65, 66, 67, 68, 69, 70, 71, 72, 73, 74, 75, 76, 77, 78, 79, 80, 81, 82, 83, 84, 85, 86, 87, 88, 89, 90, 91, 92, 93, 94, 95, 96, 97, 98, 99, 100, 125, 150, 175, 200, 225, 250, 275, 300, 325, 350, 375, 400, 425, 450, 475, 500, 550, 600, 650, 700, 750, 800, 850, 900, 950, 1000, 1250, 1500, 1750, 2000, 2250, 2500, 2750, 3000, 3500, or 4000 nucleotides long.

20 The primers for the beta4 gene, alpha6 gene, or laminin5-gamma2 subunit gene, or any of the sequences or fragments of the sequences, set forth in SEQ ID NOs:1,3,5,7,9,11, and 13, for example, typically will be used to produce an amplified DNA product that contains a desired region. In general, typically the size of the product will be such that the size can be accurately determined to within 3, or 2 or 1 nucleotides.

25 In certain embodiments this product is at least about 20, 21, 22, 23, 24, 25, 26, 27, 28, 29, 30, 31, 32, 33, 34, 35, 36, 37, 38, 39, 40, 41, 42, 43, 44, 45, 46, 47, 48, 49, 50, 51, 52, 53, 54, 55, 56, 57, 58, 59, 60, 61, 62, 63, 64, 65, 66, 67, 68, 69, 70, 71, 72, 73, 74, 75, 76, 77, 78, 79, 80, 81, 82, 83, 84, 85, 86, 87, 88, 89, 90, 91, 92, 93, 94, 95, 96, 97, 98, 99, 100, 125, 150, 175, 200, 225, 250, 275, 300, 325, 350, 375, 400, 425, 450, 475, 500, 550, 600, 650, 700, 750, 800, 850, 900, 950, 1000, 1250, 1500, 1750, 2000, 2250, 2500, 2750, 3000, 3500, or 4000 nucleotides long.

In other embodiments the product is less than or equal to about 20, 21, 22, 23, 24, 25, 26, 27, 28, 29, 30, 31, 32, 33, 34, 35, 36, 37, 38, 39, 40, 41, 42, 43, 44, 45, 46, 47, 48, 49, 50, 51, 52, 53, 54, 55, 56, 57, 58, 59, 60, 61, 62, 63, 64, 65, 66, 67, 68, 69, 70, 71, 72, 73, 74, 75, 76, 77, 78, 79, 80, 81, 82, 83, 84, 85, 86, 87, 88, 89, 90, 91, 92, 93, 94, 95, 96, 97, 98, 99, 100, 125, 150, 175,

35 200, 225, 250, 275, 300, 325, 350, 375, 400, 425, 450, 475, 500, 550, 600, 650, 700, 750, 800, 850,

900, 950, 1000, 1250, 1500, 1750, 2000, 2250, 2500, 2750, 3000, 3500, or 4000 nucleotides long.

d) Sequence similarities

It is understood that as discussed herein the use of the terms homology and identity mean the same thing as similarity. Thus, for example, if the word homology is used between two non-
5 natural sequences it is understood that this is not necessarily indicating an evolutionary relationship between these two sequences, but rather is looking at the similarity or relatedness between their nucleic acid sequences. Many of the methods for determining homology between two evolutionarily related molecules are routinely applied to any two or more nucleic acids or proteins for the purpose of measuring sequence similarity regardless of whether or not they are
10 evolutionarily related.

In general, it is understood that one way to define any known variants and derivatives or those that might arise, of the disclosed genes and proteins herein, is through defining the variants and derivatives in terms of homology to specific known sequences. This identity of particular sequences disclosed herein is also discussed elsewhere herein. In general, variants of genes and
15 proteins herein disclosed typically have at least, about 70, 71, 72, 73, 74, 75, 76, 77, 78, 79, 80, 81, 82, 83, 84, 85, 86, 87, 88, 89, 90, 91, 92, 93, 94, 95, 96, 97, 98, or 99 percent homology to the stated sequence or the native sequence. Those of skill in the art readily understand how to determine the homology of two proteins or nucleic acids, such as genes. For example, the homology can be calculated after aligning the two sequences so that the homology is at its highest
20 level.

Another way of calculating homology can be performed by published algorithms. Optimal alignment of sequences for comparison may be conducted by the local homology algorithm of Smith and Waterman Adv. Appl. Math. 2: 482 (1981), by the homology alignment algorithm of Needleman and Wunsch, J. Mol. Biol. 48: 443 (1970), by the search for similarity method of
25 Pearson and Lipman, Proc. Natl. Acad. Sci. U.S.A. 85: 2444 (1988), by computerized implementations of these algorithms (GAP, BESTFIT, FASTA, and TFASTA in the Wisconsin Genetics Software Package, Genetics Computer Group, 575 Science Dr., Madison, WI), or by inspection.

The same types of homology can be obtained for nucleic acids by for example the
30 algorithms disclosed in Zuker, M. *Science* 244:48-52, 1989, Jaeger et al. *Proc. Natl. Acad. Sci. USA* 86:7706-7710, 1989, Jaeger et al. *Methods Enzymol.* 183:281-306, 1989 which are herein incorporated by reference for at least material related to nucleic acid alignment. It is understood that any of the methods typically can be used and that in certain instances the results of these various methods may differ, but the skilled artisan understands if identity is found with at least one
35 of these methods, the sequences would be said to have the stated identity, and be disclosed herein.

For example, as used herein, a sequence recited as having a particular percent homology to another sequence refers to sequences that have the recited homology as calculated by any one or more of the calculation methods described above. For example, a first sequence has 80 percent homology, as defined herein, to a second sequence if the first sequence is calculated to have 80 percent homology to the second sequence using the Zuker calculation method even if the first sequence does not have 80 percent homology to the second sequence as calculated by any of the other calculation methods. As another example, a first sequence has 80 percent homology, as defined herein, to a second sequence if the first sequence is calculated to have 80 percent homology to the second sequence using both the Zuker calculation method and the Pearson and Lipman calculation method even if the first sequence does not have 80 percent homology to the second sequence as calculated by the Smith and Waterman calculation method, the Needleman and Wunsch calculation method, the Jaeger calculation methods, or any of the other calculation methods. As yet another example, a first sequence has 80 percent homology, as defined herein, to a second sequence if the first sequence is calculated to have 80 percent homology to the second sequence using each of calculation methods (although, in practice, the different calculation methods will often result in different calculated homology percentages).

It is also understood that functional fragments as well as antigenic fragments as well as fragments that can be used in selection protocols of the disclosed compositions are also disclosed. For example, integrins have domains that interact with the other integrins. It may be advantageous in certain embodiments to utilize just the integrin binding domain fragment of, for example, the beta4 integrin, in a selection protocol disclosed herein. By using this domain of the beta4 integrin as the selection target, for example, the selection protocol will be biased for molecules binding this domain of beta4 integrin.

e) Hybridization/selective hybridization

The term hybridization typically means a sequence driven interaction between at least two nucleic acid molecules, such as a primer or a probe and a gene. Sequence driven interaction means an interaction that occurs between two nucleotides or nucleotide analogs or nucleotide derivatives in a nucleotide specific manner. For example, G interacting with C or A interacting with T are sequence driven interactions. Typically sequence driven interactions occur on the Watson-Crick face or Hoogsteen face of the nucleotide. The hybridization of two nucleic acids is affected by a number of conditions and parameters known to those of skill in the art. For example, the salt concentrations, pH, and temperature of the reaction all affect whether two nucleic acid molecules will hybridize.

Parameters for selective hybridization between two nucleic acid molecules are well known to those of skill in the art. For example, in some embodiments selective hybridization conditions

can be defined as stringent hybridization conditions. For example, stringency of hybridization is controlled by both temperature and salt concentration of either or both of the hybridization and washing steps. For example, the conditions of hybridization to achieve selective hybridization may involve hybridization in high ionic strength solution (6X SSC or 6X SSPE) at a temperature
5 that is about 12-25°C below the T_m (the melting temperature at which half of the molecules dissociate from their hybridization partners) followed by washing at a combination of temperature and salt concentration chosen so that the washing temperature is about 5°C to 20°C below the T_m . The temperature and salt conditions are readily determined empirically in preliminary experiments in which samples of reference DNA immobilized on filters are hybridized to a labeled
10 nucleic acid of interest and then washed under conditions of different stringencies. Hybridization temperatures are typically higher for DNA-RNA and RNA-RNA hybridizations. The conditions can be used as described above to achieve stringency, or as is known in the art. (Sambrook et al., Molecular Cloning: A Laboratory Manual, 2nd Ed., Cold Spring Harbor Laboratory, Cold Spring Harbor, New York, 1989; Kunkel et al. Methods Enzymol. 1987:154:367, 1987 which is herein
15 incorporated by reference for material at least related to hybridization of nucleic acids). A preferable stringent hybridization condition for a DNA:DNA hybridization can be at about 68°C (in aqueous solution) in 6X SSC or 6X SSPE followed by washing at 68°C. Stringency of hybridization and washing, if desired, can be reduced accordingly as the degree of complementarity desired is decreased, and further, depending upon the G-C or A-T richness of any
20 area wherein variability is searched for. Likewise, stringency of hybridization and washing, if desired, can be increased accordingly as homology desired is increased, and further, depending upon the G-C or A-T richness of any area wherein high homology is desired, all as known in the art.

Another way to define selective hybridization is by looking at the amount (percentage) of
25 one of the nucleic acids bound to the other nucleic acid. For example, in some embodiments selective hybridization conditions would be when at least about, 60, 65, 70, 71, 72, 73, 74, 75, 76, 77, 78, 79, 80, 81, 82, 83, 84, 85, 86, 87, 88, 89, 90, 91, 92, 93, 94, 95, 96, 97, 98, 99, 100 percent of the limiting nucleic acid is bound to the non-limiting nucleic acid. Typically, the non-limiting primer is in for example, 10 or 100 or 1000 fold excess. This type of assay can be performed at
30 under conditions where both the limiting and non-limiting primer are for example, 10 fold or 100 fold or 1000 fold below their k_d , or where only one of the nucleic acid molecules is 10 fold or 100 fold or 1000 fold or where one or both nucleic acid molecules are above their k_d .

Another way to define selective hybridization is by looking at the percentage of primer that gets enzymatically manipulated under conditions where hybridization is required to promote
35 the desired enzymatic manipulation. For example, in some embodiments selective hybridization conditions would be when at least about, 60, 65, 70, 71, 72, 73, 74, 75, 76, 77, 78, 79, 80, 81, 82,

83, 84, 85, 86, 87, 88, 89, 90, 91, 92, 93, 94, 95, 96, 97, 98, 99, 100 percent of the primer is enzymatically manipulated under conditions which promote the enzymatic manipulation, for example if the enzymatic manipulation is DNA extension, then selective hybridization conditions would be when at least about 60, 65, 70, 71, 72, 73, 74, 75, 76, 77, 78, 79, 80, 81, 82, 83, 84, 85, 5 86, 87, 88, 89, 90, 91, 92, 93, 94, 95, 96, 97, 98, 99, 100 percent of the primer molecules are extended. Preferred conditions also include those suggested by the manufacturer or indicated in the art as being appropriate for the enzyme performing the manipulation.

Just as with homology, it is understood that there are a variety of methods herein disclosed for determining the level of hybridization between two nucleic acid molecules. It is understood 10 that these methods and conditions may provide different percentages of hybridization between two nucleic acid molecules, but unless otherwise indicated meeting the parameters of any of the methods would be sufficient. For example if 80% hybridization was required and as long as hybridization occurs within the required parameters in any one of these methods it is considered disclosed herein.

15 It is understood that those of skill in the art understand that if a composition or method meets any one of these criteria for determining hybridization either collectively or singly it is a composition or method that is disclosed herein.

f) Delivery of the compositions to cells

The disclosed compositions and methods often entail delivery of the compositions to cells. 20 For example, antisense molecules directed to alpha6 mRNA or gamma2 mRNA can be delivered to cells via any method. A number of exemplary methods are disclosed herein. It is also understood that in certain embodiments, non-nucleic acid molecules will be and can be delivered to cells, for example an antibody to beta4 integrin, alpha6 integrin or, gamma2, or a small molecule, or a peptide. Delivery of these molecules can occur by any means, and exemplary 25 compositions and methods for such delivery are disclosed herein.

There are a number of compositions and methods which can be used to deliver nucleic acids to cells, either in vitro or in vivo. These methods and compositions can largely be broken down into two classes: viral based delivery systems and non-viral based delivery systems. For example, the nucleic acids can be delivered through a number of direct delivery systems such as, 30 electroporation, lipofection, calcium phosphate precipitation, plasmids, viral vectors, viral nucleic acids, phage nucleic acids, phages, cosmids, or via transfer of genetic material in cells or carriers such as cationic liposomes. Appropriate means for transfection, including viral vectors, chemical transfectants, or physico-mechanical methods such as electroporation and direct diffusion of DNA, are described by, for example, Wolff, J. A., et al., Science, 247, 1465-1468, (1990); and Wolff, J. 35 A. Nature, 352, 815-818, (1991). Such methods are well known in the art and readily adaptable for

use with the compositions and methods described herein. In certain cases, the methods will be modified to specifically function with large DNA molecules. Further, these methods can be used to target certain diseases and cell populations by using the targeting characteristics of the carrier.

(1) Nucleic acid based delivery systems

5 Transfer vectors can be any nucleotide construction used to deliver genes into cells (e.g., a plasmid), or as part of a general strategy to deliver genes, e.g., as part of recombinant retrovirus or adenovirus (Ram et al. Cancer Res. 53:83-88, (1993)).

As used herein, plasmid or viral vectors are agents that transport the disclosed nucleic acids, such as gamma2 antisense producing molecules into the cell without degradation and
10 include a promoter yielding expression of the gene in the cells into which it is delivered. In some embodiments the delivery vectors are derived from either a virus or a retrovirus. Viral vectors are, for example, Adenovirus, Adeno-associated virus, Herpes virus, Vaccinia virus, Polio virus, AIDS virus, neuronal trophic virus, Sindbis and other RNA viruses, including these viruses with the HIV backbone. Also preferred are any viral families which share the properties of these viruses which
15 make them suitable for use as vectors. Retroviruses include Murine Moloney Leukemia virus, MMLV, and retroviruses that express the desirable properties of MMLV as a vector. Retroviral vectors are able to carry a larger genetic payload, i.e., a transgene or marker gene, than other viral vectors, and for this reason are a commonly used vector. However, other than Lentivirus vectors, they are typically not as useful in non-proliferating cells. Adenovirus vectors are relatively stable
20 and easy to work with, have high titers, and can be delivered in aerosol formulation, and can transfect non-dividing cells. Pox viral vectors are large and have several sites for inserting genes, they are thermostable and can be stored at room temperature. A preferred embodiment is a viral vector which has been engineered so as to suppress the immune response of the host organism, elicited by the viral antigens. Preferred vectors of this type will carry coding regions for
25 Interleukin 8 or 10.

Viral vectors can have higher transaction (ability to introduce genes) abilities than chemical or physical methods to introduce genes into cells. Typically, viral vectors contain, nonstructural early genes, structural late genes, an RNA polymerase III transcript, inverted terminal repeats necessary for replication and encapsidation, and promoters to control the
30 transcription and replication of the viral genome. When engineered as vectors, viruses typically have one or more of the early genes removed and a gene or gene/promoter cassette is inserted into the viral genome in place of the removed viral DNA. Constructs of this type can carry up to about 8 kb of foreign genetic material. The necessary functions of the removed early genes are typically supplied by cell lines which have been engineered to express the gene products of the early genes
35 in trans.

(a) Retroviral Vectors

A retrovirus is an animal virus belonging to the virus family of Retroviridae, including any types, subfamilies, genus, or tropisms. Retroviral vectors, in general, are described by Verma, I.M., Retroviral vectors for gene transfer. In Microbiology-1985, American Society for
5 Microbiology, pp. 229-232, Washington, (1985), which is incorporated by reference herein. Examples of methods for using retroviral vectors for gene therapy are described in U.S. Patent Nos. 4,868,116 and 4,980,286; PCT applications WO 90/02806 and WO 89/07136; and Mulligan, (Science 260:926-932 (1993)); the teachings of which are incorporated herein by reference.

A retrovirus is essentially a package which has packed into it nucleic acid cargo. The
10 nucleic acid cargo carries with it a packaging signal, which ensures that the replicated daughter molecules will be efficiently packaged within the package coat. In addition to the package signal, there are a number of molecules which are needed in cis, for the replication, and packaging of the replicated virus. Typically a retroviral genome, contains the gag, pol, and env genes which are involved in the making of the protein coat. It is the gag, pol, and env genes which are typically
15 replaced by the foreign DNA that it is to be transferred to the target cell. Retrovirus vectors typically contain a packaging signal for incorporation into the package coat, a sequence which signals the start of the gag transcription unit, elements necessary for reverse transcription, including a primer binding site to bind the tRNA primer of reverse transcription, terminal repeat sequences that guide the switch of RNA strands during DNA synthesis, a purine rich sequence 5' to
20 the 3' LTR that serve as the priming site for the synthesis of the second strand of DNA synthesis, and specific sequences near the ends of the LTRs that enable the insertion of the DNA state of the retrovirus to insert into the host genome. The removal of the gag, pol, and env genes allows for about 8 kb of foreign sequence to be inserted into the viral genome, become reverse transcribed, and upon replication be packaged into a new retroviral particle. This amount of nucleic acid is
25 sufficient for the delivery of a one to many genes depending on the size of each transcript. It is preferable to include either positive or negative selectable markers along with other genes in the insert.

Since the replication machinery and packaging proteins in most retroviral vectors have been removed (gag, pol, and env), the vectors are typically generated by placing them into a
30 packaging cell line. A packaging cell line is a cell line which has been transfected or transformed with a retrovirus that contains the replication and packaging machinery, but lacks any packaging signal. When the vector carrying the DNA of choice is transfected into these cell lines, the vector containing the gene of interest is replicated and packaged into new retroviral particles, by the machinery provided in cis by the helper cell. The genomes for the machinery are not packaged
35 because they lack the necessary signals.

(b) Adenoviral Vectors

The construction of replication-defective adenoviruses has been described (Berkner et al., J. Virology 61:1213-1220 (1987); Massie et al., Mol. Cell. Biol. 6:2872-2883 (1986); Haj-Ahmad et al., J. Virology 57:267-274 (1986); Davidson et al., J. Virology 61:1226-1239 (1987);

5 Zhang "Generation and identification of recombinant adenovirus by liposome-mediated transfection and PCR analysis" BioTechniques 15:868-872 (1993)). The benefit of the use of these viruses as vectors is that they are limited in the extent to which they can spread to other cell types, since they can replicate within an initial infected cell, but are unable to form new infectious viral particles. Recombinant adenoviruses have been shown to achieve high efficiency gene

10 transfer after direct, in vivo delivery to airway epithelium, hepatocytes, vascular endothelium, CNS parenchyma and a number of other tissue sites (Morsy, J. Clin. Invest. 92:1580-1586 (1993); Kirshenbaum, J. Clin. Invest. 92:381-387 (1993); Roessler, J. Clin. Invest. 92:1085-1092 (1993); Moullier, Nature Genetics 4:154-159 (1993); La Salle, Science 259:988-990 (1993); Gomez-Foix, J. Biol. Chem. 267:25129-25134 (1992); Rich, Human Gene Therapy 4:461-476

15 (1993); Zabner, Nature Genetics 6:75-83 (1994); Guzman, Circulation Research 73:1201-1207 (1993); Bout, Human Gene Therapy 5:3-10 (1994); Zabner, Cell 75:207-216 (1993); Caillaud, Eur. J. Neuroscience 5:1287-1291 (1993); and Ragot, J. Gen. Virology 74:501-507 (1993)). Recombinant adenoviruses achieve gene transduction by binding to specific cell surface receptors, after which the virus is internalized by receptor-mediated endocytosis, in the same manner as wild

20 type or replication-defective adenovirus (Chardonnet and Dales, Virology 40:462-477 (1970); Brown and Burlingham, J. Virology 12:386-396 (1973); Svensson and Persson, J. Virology 55:442-449 (1985); Seth, et al., J. Virol. 51:650-655 (1984); Seth, et al., Mol. Cell. Biol. 4:1528-1533 (1984); Varga et al., J. Virology 65:6061-6070 (1991); Wickham et al., Cell 73:309-319 (1993)).

25 A viral vector can be one based on an adenovirus which has had the E1 gene removed and these virions are generated in a cell line such as the human 293 cell line. In another preferred embodiment both the E1 and E3 genes are removed from the adenovirus genome.

(c) Adeno-associated viral vectors

Another type of viral vector is based on an adeno-associated virus (AAV). This defective

30 parvovirus is a preferred vector because it can infect many cell types and is nonpathogenic to humans. AAV type vectors can transport about 4 to 5 kb and wild type AAV is known to stably insert into chromosome 19. Vectors which contain this site specific integration property are preferred. An especially preferred embodiment of this type of vector is the P4.1 C vector produced by Avigen, San Francisco, CA, which can contain the herpes simplex virus thymidine kinase gene,

35 HSV-tk, and/or a marker gene, such as the gene encoding the green fluorescent protein, GFP.

In another type of AAV virus, the AAV contains a pair of inverted terminal repeats (ITRs) which flank at least one cassette containing a promoter which directs cell-specific expression operably linked to a heterologous gene. Heterologous in this context refers to any nucleotide sequence or gene which is not native to the AAV or B19 parvovirus.

5 Typically the AAV and B19 coding regions have been deleted, resulting in a safe, noncytotoxic vector. The AAV ITRs, or modifications thereof, confer infectivity and site-specific integration, but not cytotoxicity, and the promoter directs cell-specific expression. United states Patent No. 6,261,834 is herein incorporated by reference for material related to the AAV vector.

The vectors of the present invention thus provide DNA molecules which are capable of
10 integration into a mammalian chromosome without substantial toxicity.

The inserted genes in viral and retroviral usually contain promoters, and/or enhancers to help control the expression of the desired gene product. A promoter is generally a sequence or sequences of DNA that function when in a relatively fixed location in regard to the transcription start site. A promoter contains core elements required for basic interaction of RNA polymerase
15 and transcription factors, and may contain upstream elements and response elements.

(d) Large payload viral vectors

Molecular genetic experiments with large human herpesviruses have provided a means whereby large heterologous DNA fragments can be cloned, propagated and established in cells permissive for infection with herpesviruses (Sun et al., Nature genetics 8: 33-41, 1994; Cotter and
20 Robertson, Curr Opin Mol Ther 5: 633-644, 1999). These large DNA viruses (herpes simplex virus (HSV) and Epstein-Barr virus (EBV), have the potential to deliver fragments of human heterologous DNA > 150 kb to specific cells. EBV recombinants can maintain large pieces of DNA in the infected B-cells as episomal DNA. Individual clones carried human genomic inserts up to 330 kb appeared genetically stable. The maintenance of these episomes requires a specific
25 EBV nuclear protein, EBNA1, constitutively expressed during infection with EBV. Additionally, these vectors can be used for transfection, where large amounts of protein can be generated transiently in vitro. Herpesvirus amplicon systems are also being used to package pieces of DNA > 220 kb and to infect cells that can stably maintain DNA as episomes.

Other useful systems include, for example, replicating and host-restricted non-replicating
30 vaccinia virus vectors.

(2) Non-nucleic acid based systems

The disclosed compositions can be delivered to the target cells in a variety of ways. For example, the compositions can be delivered through electroporation, or through lipofection, or through calcium phosphate precipitation. The delivery mechanism chosen will depend in part on

the type of cell targeted and whether the delivery is occurring for example *in vivo* or *in vitro*.

Thus, the compositions can comprise, in addition to the disclosed nucleic acids and proteins or vectors for example, lipids such as liposomes, such as cationic liposomes (e.g., DOTMA, DOPE, DC-cholesterol) or anionic liposomes. Liposomes can further comprise proteins
5 to facilitate targeting a particular cell, if desired. Administration of a composition comprising a compound and a cationic liposome can be administered to the blood afferent to a target organ or inhaled into the respiratory tract to target cells of the respiratory tract. Regarding liposomes, see, e.g., Brigham et al. *Am. J. Resp. Cell. Mol. Biol.* 1:95-100 (1989); Felgner et al. *Proc. Natl. Acad. Sci USA* 84:7413-7417 (1987); U.S. Pat. No.4,897,355. Furthermore, the compound can
10 be administered as a component of a microcapsule that can be targeted to specific cell types, such as macrophages, or where the diffusion of the compound or delivery of the compound from the microcapsule is designed for a specific rate or dosage.

In the methods described above which include the administration and uptake of exogenous DNA into the cells of a subject (i.e., gene transduction or transfection), delivery of the
15 compositions to cells can be via a variety of mechanisms. As one example, delivery can be via a liposome, using commercially available liposome preparations such as LIPOFECTIN, LIPOFECTAMINE (GIBCO-BRL, Inc., Gaithersburg, MD), SUPERFECT (Qiagen, Inc. Hilden, Germany) and TRANSFECTAM (Promega Biotec, Inc., Madison, WI), as well as other liposomes developed according to procedures standard in the art. In addition, the nucleic acid or vector of
20 this invention can be delivered *in vivo* by electroporation, the technology for which is available from Genetronics, Inc. (San Diego, CA) as well as by means of a SONOPORATION machine (ImaRx Pharmaceutical Corp., Tucson, AZ).

The materials may be in solution or suspension (for example, incorporated into microparticles, liposomes, or cells). These may be targeted to a particular cell type via antibodies,
25 receptors, or receptor ligands. The following references are examples of the use of this technology to target specific proteins to tumor tissue (Senter, et al., Bioconjugate Chem., 2:447-451, (1991); Bagshawe, K.D., Br. J. Cancer, 60:275-281, (1989); Bagshawe, et al., Br. J. Cancer, 58:700-703, (1988); Senter, et al., Bioconjugate Chem., 4:3-9, (1993); Battelli, et al., Cancer Immunol. Immunother., 35:421-425, (1992); Pietersz and McKenzie, Immunolog. Reviews, 129:57-80, (1992); and Roffler, et al., Biochem. Pharmacol., 42:2062-2065, (1991)). These techniques can be
30 used for a variety of other specific cell types. Vehicles, among others, include "stealth" and other antibody conjugated liposomes (including lipid mediated drug targeting to colonic carcinoma), receptor mediated targeting of DNA through cell specific ligands, lymphocyte directed tumor targeting, and highly specific therapeutic retroviral targeting of murine glioma cells *in vivo*. The
35 following references are examples of the use of this technology to target specific proteins to tumor

tissue (Hughes et al., Cancer Research, 49:6214-6220, (1989); and Litzinger and Huang, Biochimica et Biophysica Acta, 1104:179-187, (1992)). In general, receptors are involved in pathways of endocytosis, either constitutive or ligand induced. These receptors cluster in clathrin-coated pits, enter the cell via clathrin-coated vesicles, pass through an acidified endosome in which
5 the receptors are sorted, and then either recycle to the cell surface, become stored intracellularly, or are degraded in lysosomes. The internalization pathways serve a variety of functions, such as nutrient uptake, removal of activated proteins, clearance of macromolecules, opportunistic entry of viruses and toxins, dissociation and degradation of ligand, and receptor-level regulation. Many receptors follow more than one intracellular pathway, depending on the cell type, receptor
10 concentration, type of ligand, ligand valency, and ligand concentration. Molecular and cellular mechanisms of receptor-mediated endocytosis has been reviewed (Brown and Greene, DNA and Cell Biology 10:6, 399-409 (1991)).

Nucleic acids that are delivered to cells which are to be integrated into the host cell genome, typically contain integration sequences. These sequences are often viral related
15 sequences, particularly when viral based systems are used. These viral integration systems can also be incorporated into nucleic acids which are to be delivered using a non-nucleic acid based system of deliver, such as a liposome, so that the nucleic acid contained in the delivery system can be come integrated into the host genome.

Other general techniques for integration into the host genome include, for example,
20 systems designed to promote homologous recombination with the host genome. These systems typically rely on sequence flanking the nucleic acid to be expressed that has enough homology with a target sequence within the host cell genome that recombination between the vector nucleic acid and the target nucleic acid takes place, causing the delivered nucleic acid to be integrated into the host genome. These systems and the methods necessary to promote homologous
25 recombination are known to those of skill in the art.

(3) In vivo/ex vivo

As described above, the compositions can be administered in a pharmaceutically acceptable carrier and can be delivered to the subject's cells *in vivo* and/or *ex vivo* by a variety of mechanisms well known in the art (e.g., uptake of naked DNA, liposome fusion, intramuscular
30 injection of DNA via a gene gun, endocytosis and the like).

If *ex vivo* methods are employed, cells or tissues can be removed and maintained outside the body according to standard protocols well known in the art. The compositions can be introduced into the cells via any gene transfer mechanism, such as, for example, calcium phosphate mediated gene delivery, electroporation, microinjection or proteoliposomes. The
35 transduced cells can then be infused (e.g., in a pharmaceutically acceptable carrier) or

homotopically transplanted back into the subject per standard methods for the cell or tissue type. Standard methods are known for transplantation or infusion of various cells into a subject.

It is understood that in certain embodiments, constructs which produce an integrin signal transduction inhibitor are driven by inducible promoters, rather than constitutive promoters.

- 5 inducible systems provide certain advantages, to the expression of the disclosed constructs. Any inducible system can be used. Also disclosed are cells containing the inducible systems, described herein, and in the Examples. These cells, can be used as model systems in a wide variety of assays, as well as in vivo settings.

g) Expression systems

- 10 The nucleic acids that are delivered to cells typically contain expression controlling systems. For example, the inserted genes in viral and retroviral systems usually contain promoters, and/or enhancers to help control the expression of the desired gene product. A promoter is generally a sequence or sequences of DNA that function when in a relatively fixed location in regard to the transcription start site. A promoter contains core elements required for basic
15 interaction of RNA polymerase and transcription factors, and may contain upstream elements and response elements.

(1) Viral Promoters and Enhancers

- Preferred promoters controlling transcription from vectors in mammalian host cells may be obtained from various sources, for example, the genomes of viruses such as: polyoma, Simian
20 Virus 40 (SV40), adenovirus, retroviruses, hepatitis-B virus and most preferably cytomegalovirus, or from heterologous mammalian promoters, e.g. beta actin promoter. The early and late promoters of the SV40 virus are conveniently obtained as an SV40 restriction fragment which also contains the SV40 viral origin of replication (Fiers et al., Nature, 273: 113 (1978)). The immediate early promoter of the human cytomegalovirus is conveniently obtained as a HindIII E
25 restriction fragment (Greenway, P.J. et al., Gene 18: 355-360 (1982)). Of course, promoters from the host cell or related species also are useful herein.

- Enhancer generally refers to a sequence of DNA that functions at no fixed distance from the transcription start site and can be either 5' (Laimins, L. et al., Proc. Natl. Acad. Sci. 78: 993 (1981)) or 3' (Lusky, M.L., et al., Mol. Cell Bio. 3: 1108 (1983)) to the transcription unit.
30 Furthermore, enhancers can be within an intron (Banerji, J.L. et al., Cell 33: 729 (1983)) as well as within the coding sequence itself (Osborne, T.F., et al., Mol. Cell Bio. 4: 1293 (1984)). They are usually between 10 and 300 bp in length, and they function in cis. Enhancers function to increase transcription from nearby promoters. Enhancers also often contain response elements that mediate the regulation of transcription. Promoters can also contain response elements that mediate
35 the regulation of transcription. Enhancers often determine the regulation of expression of a gene.

While many enhancer sequences are now known from mammalian genes (globin, elastase, albumin, α -fetoprotein and insulin), typically one will use an enhancer from a eukaryotic cell virus for general expression. Preferred examples are the SV40 enhancer on the late side of the replication origin (bp 100-270), the cytomegalovirus early promoter enhancer, the polyoma
5 enhancer on the late side of the replication origin, and adenovirus enhancers.

The promoter and/or enhancer may be specifically activated either by light or specific chemical events which trigger their function. Systems can be regulated by reagents such as tetracycline and dexamethasone. There are also ways to enhance viral vector gene expression by exposure to irradiation, such as gamma irradiation, or alkylating chemotherapy drugs.

10 In certain embodiments the promoter and/or enhancer region can act as a constitutive promoter and/or enhancer to maximize expression of the region of the transcription unit to be transcribed. In certain constructs the promoter and/or enhancer region be active in all eukaryotic cell types, even if it is only expressed in a particular type of cell at a particular time. A preferred promoter of this type is the CMV promoter (650 bases). Other preferred promoters are SV40
15 promoters, cytomegalovirus (full length promoter), and retroviral vector LTF.

It has been shown that all specific regulatory elements can be cloned and used to construct expression vectors that are selectively expressed in specific cell types such as melanoma cells. The glial fibrillary acetic protein (GFAP) promoter has been used to selectively express genes in cells of glial origin.

20 Expression vectors used in eukaryotic host cells (yeast, fungi, insect, plant, animal, human or nucleated cells) may also contain sequences necessary for the termination of transcription which may affect mRNA expression. These regions are transcribed as polyadenylated segments in the untranslated portion of the mRNA encoding tissue factor protein. The 3' untranslated regions also include transcription termination sites. It is preferred that the transcription unit also contain a
25 polyadenylation region. One benefit of this region is that it increases the likelihood that the transcribed unit will be processed and transported like mRNA. The identification and use of polyadenylation signals in expression constructs is well established. It is preferred that homologous polyadenylation signals be used in the transgene constructs. In certain transcription units, the polyadenylation region is derived from the SV40 early polyadenylation signal and
30 consists of about 400 bases. It is also preferred that the transcribed units contain other standard sequences alone or in combination with the above sequences improve expression from, or stability of, the construct.

(2) Markers

The viral vectors can include nucleic acid sequence encoding a marker product. This
35 marker product is used to determine if the gene has been delivered to the cell and once delivered is

being expressed. Preferred marker genes are the *E. Coli* lacZ gene, which encodes β -galactosidase, and green fluorescent protein.

In some embodiments the marker may be a selectable marker. Examples of suitable selectable markers for mammalian cells are dihydrofolate reductase (DHFR), thymidine kinase, 5 neomycin, neomycin analog G418, hydromycin, and puromycin. When such selectable markers are successfully transferred into a mammalian host cell, the transformed mammalian host cell can survive if placed under selective pressure. There are two widely used distinct categories of selective regimes. The first category is based on a cell's metabolism and the use of a mutant cell line which lacks the ability to grow independent of a supplemented media. Two examples are: 10 CHO DHFR- cells and mouse LTK- cells. These cells lack the ability to grow without the addition of such nutrients as thymidine or hypoxanthine. Because these cells lack certain genes necessary for a complete nucleotide synthesis pathway, they cannot survive unless the missing nucleotides are provided in a supplemented media. An alternative to supplementing the media is to introduce an intact DHFR or TK gene into cells lacking the respective genes, thus altering their 15 growth requirements. Individual cells which were not transformed with the DHFR or TK gene will not be capable of survival in non-supplemented media.

The second category is dominant selection which refers to a selection scheme used in any cell type and does not require the use of a mutant cell line. These schemes typically use a drug to arrest growth of a host cell. Those cells which have a novel gene would express a protein 20 conveying drug resistance and would survive the selection. Examples of such dominant selection use the drugs neomycin, (Southern P. and Berg, P., J. Molec. Appl. Genet. 1: 327 (1982)), mycophenolic acid, (Mulligan, R.C. and Berg, P. Science 209: 1422 (1980)) or hygromycin, (Sugden, B. et al., Mol. Cell. Biol. 5: 410-413 (1985)). The three examples employ bacterial genes under eukaryotic control to convey resistance to the appropriate drug G418 or neomycin 25 (geneticin), xgpt (mycophenolic acid) or hygromycin, respectively. Others include the neomycin analog G418 and puromycin.

h) Peptides

(1) Protein variants

As discussed herein there are numerous variants of the beta4 integrin protein, alpha6 30 integrin protein, and gamma2 laminin5 protein, for example, that are known and herein contemplated. In addition, to the known functional homologue variants there are derivatives of the beta4, alpha6, and gamma2, and other disclosed proteins which also function in the disclosed methods and compositions. Protein variants and derivatives are well understood to those of skill in the art and in can involve amino acid sequence modifications. For example, amino acid sequence 35 modifications typically fall into one or more of three classes: substitutional, insertional or

deletional variants. Insertions include amino and/or carboxyl terminal fusions as well as intrasequence insertions of single or multiple amino acid residues. Insertions ordinarily will be smaller insertions than those of amino or carboxyl terminal fusions, for example, on the order of one to four residues. Immunogenic fusion protein derivatives are made by fusing a polypeptide sufficiently large to confer immunogenicity to the target sequence by cross-linking in vitro or by recombinant cell culture transformed with DNA encoding the fusion. Deletions are characterized by the removal of one or more amino acid residues from the protein sequence. Typically, no more than about from 2 to 6 residues are deleted at any one site within the protein molecule. These variants ordinarily are prepared by site specific mutagenesis of nucleotides in the DNA encoding the protein, thereby producing DNA encoding the variant, and thereafter expressing the DNA in recombinant cell culture. Techniques for making substitution mutations at predetermined sites in DNA having a known sequence are well known, for example M13 primer mutagenesis and PCR mutagenesis. Amino acid substitutions are typically of single residues, but can occur at a number of different locations at once; insertions usually will be on the order of about from 1 to 10 amino acid residues; and deletions will range about from 1 to 30 residues. Deletions or insertions preferably are made in adjacent pairs, i.e. a deletion of 2 residues or insertion of 2 residues. Substitutions, deletions, insertions or any combination thereof may be combined to arrive at a final construct. The mutations must not place the sequence out of reading frame and preferably will not create complementary regions that could produce secondary mRNA structure. Substitutional variants are those in which at least one residue has been removed and a different residue inserted in its place. Such substitutions generally are made in accordance with the following Tables 1 and 2 and are referred to as conservative substitutions.

TABLE 1: Amino Acid Abbreviations

Amino Acid	Abbreviations	
alanine	Ala	A
allosoleucine	AlIle	
arginine	Arg	R
asparagine	Asn	N
aspartic acid	Asp	D
cysteine	Cys	C
glutamic acid	Glu	E
glutamine	Gln	K
glycine	Gly	G
histidine	His	H
isoleucine	Ile	I
leucine	Leu	L
lysine	Lys	K
phenylalanine	Phe	F
proline	Pro	P
pyroglutamic acid	pGlu	
serine	Ser	S
threonine	Thr	T

tyrosine	Tyr	Y	
tryptophan		Trp	W
valine		Val	V

5	TABLE 2:	Amino Acid Substitutions
	Original Residue	Exemplary Conservative Substitutions, others are known in the art.
	Ala	ser
	Arg	lys, gln
	Asn	gln; his
10	Asp	glu
	Cys	ser
	Gln	asn, lys
	Glu	asp
	Gly	pro
15	His	asn;gln
	Ile	leu; val
	Leu	ile; val
	Lys	arg; gln;
	Met	Leu; ile
20	Phe	met; leu; tyr
	Ser	thr
	Thr	ser
	Trp	tyr
	Tyr	trp; phe
25	Val	ile; leu

Substantial changes in function or immunological identity are made by selecting substitutions that are less conservative than those in Table 2, i.e., selecting residues that differ more significantly in their effect on maintaining (a) the structure of the polypeptide backbone in the area of the substitution, for example as a sheet or helical conformation, (b) the charge or hydrophobicity of the molecule at the target site or (c) the bulk of the side chain. The substitutions which in general are expected to produce the greatest changes in the protein properties will be those in which (a) a hydrophilic residue, e.g. seryl or threonyl, is substituted for (or by) a hydrophobic residue, e.g. leucyl, isoleucyl, phenylalanyl, valyl or alanyl; (b) a cysteine or proline is substituted for (or by) any other residue; (c) a residue having an electropositive side chain, e.g., lysyl, arginyl, or histidyl, is substituted for (or by) an electronegative residue, e.g., glutamyl or aspartyl; or (d) a residue having a bulky side chain, e.g., phenylalanine, is substituted for (or by) one not having a side chain, e.g., glycine, in this case, (e) by increasing the number of sites for sulfation and/or glycosylation.

For example, the replacement of one amino acid residue with another that is biologically and/or chemically similar is known to those skilled in the art as a conservative substitution. For example, a conservative substitution would be replacing one hydrophobic residue for another, or one polar residue for another. The substitutions include combinations such as, for example, Gly, Ala; Val, Ile, Leu; Asp, Glu; Asn, Gln; Ser, Thr; Lys, Arg; and Phe, Tyr. Such conservatively substituted variations of each explicitly disclosed sequence are included within the mosaic

polypeptides provided herein.

Substitutional or deletional mutagenesis can be employed to insert sites for N-glycosylation (Asn-X-Thr/Ser) or O-glycosylation (Ser or Thr). Deletions of cysteine or other labile residues also may be desirable. Deletions or substitutions of potential proteolysis sites, e.g. Arg, is accomplished for example by deleting one of the basic residues or substituting one by glutaminyl or histidyl residues.

Certain post-translational derivatizations are the result of the action of recombinant host cells on the expressed polypeptide. Glutaminyl and asparaginyl residues are frequently post-translationally deamidated to the corresponding glutamyl and aspartyl residues. Alternatively, these residues are deamidated under mildly acidic conditions. Other post-translational modifications include hydroxylation of proline and lysine, phosphorylation of hydroxyl groups of seryl or threonyl residues, methylation of the o-amino groups of lysine, arginine, and histidine side chains (T.E. Creighton, *Proteins: Structure and Molecular Properties*, W. H. Freeman & Co., San Francisco pp 79-86 [1983]), acetylation of the N-terminal amine and, in some instances, amidation of the C-terminal carboxyl.

It is understood that one way to define the variants and derivatives of the disclosed proteins herein is through defining the variants and derivatives in terms of homology/identity to specific known sequences. For example, SEQ ID NO:5 sets forth a particular sequence of beta4 integrin cDNA and SEQ ID NO:6 sets forth a particular sequence of a beta4 integrin protein. Specifically disclosed are variants of these and other proteins herein disclosed which have at least, 70% or 75% or 80% or 85% or 90% or 95% homology to the stated sequence. Those of skill in the art readily understand how to determine the homology of two proteins. For example, the homology can be calculated after aligning the two sequences so that the homology is at its highest level.

Another way of calculating homology can be performed by published algorithms. Optimal alignment of sequences for comparison may be conducted by the local homology algorithm of Smith and Waterman *Adv. Appl. Math.* 2: 482 (1981), by the homology alignment algorithm of Needleman and Wunsch, *J. Mol. Biol.* 48: 443 (1970), by the search for similarity method of Pearson and Lipman, *Proc. Natl. Acad. Sci. U.S.A.* 85: 2444 (1988), by computerized implementations of these algorithms (GAP, BESTFIT, FASTA, and TFASTA in the Wisconsin Genetics Software Package, Genetics Computer Group, 575 Science Dr., Madison, WI), or by inspection.

The same types of homology can be obtained for nucleic acids by for example the algorithms disclosed in Zuker, M. *Science* 244:48-52, 1989, Jaeger et al. *Proc. Natl. Acad. Sci. USA* 86:7706-7710, 1989, Jaeger et al. *Methods Enzymol.* 183:281-306, 1989 which are herein

incorporated by reference for at least material related to nucleic acid alignment.

It is understood that the description of conservative mutations and homology can be combined together in any combination, such as embodiments that have at least 70% homology to a particular sequence wherein the variants are conservative mutations.

5 As this specification discusses various proteins and protein sequences it is understood that the nucleic acids that can encode those protein sequences are also disclosed. This would include all degenerate sequences related to a specific protein sequence, i.e. all nucleic acids having a sequence that encodes one particular protein sequence as well as all nucleic acids, including degenerate nucleic acids, encoding the disclosed variants and derivatives of the protein sequences.

10 Thus, while each particular nucleic acid sequence may not be written out herein, it is understood that each and every sequence is in fact disclosed and described herein through the disclosed protein sequence. For example, one of the many nucleic acid sequences that can encode the protein sequence set forth in SEQ ID NO:6 is set forth in SEQ ID NO:5. Another nucleic acid sequence that encodes the same protein sequence set forth in SEQ ID NO:6 is set forth in SEQ ID

15 NO:16. In addition, for example, a disclosed conservative derivative of SEQ ID NO:2 is shown in SEQ ID NO: 17, where the valine (V) at position 34 is changed to a isoleucine (I). It is understood that for this mutation all of the nucleic acid sequences that encode this particular derivative of the beta4 integrin are also disclosed including for example SEQ ID NO:18 and SEQ ID NO:19 which set forth two of the degenerate nucleic acid sequences that encode the particular polypeptide set

20 forth in SEQ ID NO:17. It is also understood that while no amino acid sequence indicates what particular DNA sequence encodes that protein within an organism, where particular variants of a disclosed protein are disclosed herein, the known nucleic acid sequence that encodes that protein in the particular organism from which that protein arises is also known and herein disclosed and described.

25 i) Pharmaceutical carriers/Delivery of pharmaceutical products

As described above, the compositions, for example, compositions that inhibit alpha6 function, beta4 function, or gamma2 function can also be administered *in vivo* in a pharmaceutically acceptable carrier. By "pharmaceutically acceptable" is meant a material that is not biologically or otherwise undesirable, i.e., the material may be administered to a subject, along

30 with the nucleic acid or vector, without causing any undesirable biological effects or interacting in a deleterious manner with any of the other components of the pharmaceutical composition in which it is contained. The carrier would naturally be selected to minimize any degradation of the active ingredient and to minimize any adverse side effects in the subject, as would be well known to one of skill in the art.

35 The compositions may be administered orally, parenterally (e.g., intravenously), by

intramuscular injection, by intraperitoneal injection, transdermally, extracorporeally, topically or the like, although topical intranasal administration or administration by inhalant is typically preferred. As used herein, "topical intranasal administration" means delivery of the compositions into the nose and nasal passages through one or both of the nares and can comprise delivery by a spraying mechanism or droplet mechanism, or through aerosolization of the nucleic acid or vector. The latter may be effective when a large number of animals is to be treated simultaneously. Administration of the compositions by inhalant can be through the nose or mouth via delivery by a spraying or droplet mechanism. Delivery can also be directly to any area of the respiratory system (e.g., lungs) via intubation. The exact amount of the compositions required will vary from subject to subject, depending on the species, age, weight and general condition of the subject, the severity of the allergic disorder being treated, the particular nucleic acid or vector used, its mode of administration and the like. Thus, it is not possible to specify an exact amount for every composition. However, an appropriate amount can be determined by one of ordinary skill in the art using only routine experimentation given the teachings herein.

Parenteral administration of the composition, if used, is generally characterized by injection. Injectables can be prepared in conventional forms, either as liquid solutions or suspensions, solid forms suitable for solution or suspension in liquid prior to injection, or as emulsions. A more recently revised approach for parenteral administration involves use of a slow release or sustained release system such that a constant dosage is maintained. See, e.g., U.S. Patent No. 3,610,795, which is incorporated by reference herein.

The materials may be in solution, suspension (for example, incorporated into microparticles, liposomes, or cells). These may be targeted to a particular cell type via antibodies, receptors, or receptor ligands. The following references are examples of the use of this technology to target specific proteins to tumor tissue (Senter, et al., Bioconjugate Chem., 2:447-451, (1991); Bagshawe, K.D., Br. J. Cancer, 60:275-281, (1989); Bagshawe, et al., Br. J. Cancer, 58:700-703, (1988); Senter, et al., Bioconjugate Chem., 4:3-9, (1993); Battelli, et al., Cancer Immunol. Immunother., 35:421-425, (1992); Pietersz and McKenzie, Immunolog. Reviews, 129:57-80, (1992); and Roffler, et al., Biochem. Pharmacol., 42:2062-2065, (1991)). Vehicles such as "stealth" and other antibody conjugated liposomes (including lipid mediated drug targeting to colonic carcinoma), receptor mediated targeting of DNA through cell specific ligands, lymphocyte directed tumor targeting, and highly specific therapeutic retroviral targeting of murine glioma cells *in vivo*. The following references are examples of the use of this technology to target specific proteins to tumor tissue (Hughes et al., Cancer Research, 49:6214-6220, (1989); and Litzinger and Huang, Biochimica et Biophysica Acta, 1104:179-187, (1992)). In general, receptors are involved in pathways of endocytosis, either constitutive or ligand induced. These receptors cluster in clathrin-coated pits, enter the cell via clathrin-coated vesicles, pass through an acidified endosome

in which the receptors are sorted, and then either recycle to the cell surface, become stored intracellularly, or are degraded in lysosomes. The internalization pathways serve a variety of functions, such as nutrient uptake, removal of activated proteins, clearance of macromolecules, opportunistic entry of viruses and toxins, dissociation and degradation of ligand, and receptor-level
5 regulation. Many receptors follow more than one intracellular pathway, depending on the cell type, receptor concentration, type of ligand, ligand valency, and ligand concentration. Molecular and cellular mechanisms of receptor-mediated endocytosis has been reviewed (Brown and Greene, DNA and Cell Biology 10:6, 399-409 (1991)).

(1) Pharmaceutically Acceptable Carriers

10 The compositions, including antibodies, can be used therapeutically in combination with a pharmaceutically acceptable carrier.

Pharmaceutical carriers are known to those skilled in the art. These most typically would be standard carriers for administration of drugs to humans, including solutions such as sterile water, saline, and buffered solutions at physiological pH. The compositions can be administered
15 intramuscularly or subcutaneously. Other compounds will be administered according to standard procedures used by those skilled in the art.

Pharmaceutical compositions may include carriers, thickeners, diluents, buffers, preservatives, surface active agents and the like in addition to the molecule of choice.

Pharmaceutical compositions may also include one or more active ingredients such as antimicrobial
20 agents, anti-inflammatory agents, anesthetics, and the like.

The pharmaceutical composition may be administered in a number of ways depending on whether local or systemic treatment is desired, and on the area to be treated. Administration may be topically (including ophthalmically, vaginally, rectally, intranasally), orally, by inhalation, or parenterally, for example by intravenous drip, subcutaneous, intraperitoneal or intramuscular
25 injection. The disclosed antibodies can be administered intravenously, intraperitoneally, intramuscularly, subcutaneously, intracavity, or transdermally.

Preparations for parenteral administration include sterile aqueous or non-aqueous solutions, suspensions, and emulsions. Examples of non-aqueous solvents are propylene glycol, polyethylene glycol, vegetable oils such as olive oil, and injectable organic esters such as ethyl
30 oleate. Aqueous carriers include water, alcoholic/aqueous solutions, emulsions or suspensions, including saline and buffered media. Parenteral vehicles include sodium chloride solution, Ringer's dextrose, dextrose and sodium chloride, lactated Ringer's, or fixed oils. Intravenous vehicles include fluid and nutrient replenishers, electrolyte replenishers (such as those based on Ringer's dextrose), and the like. Preservatives and other additives may also be present such as, for

example, antimicrobials, anti-oxidants, chelating agents, and inert gases and the like.

Formulations for topical administration may include ointments, lotions, creams, gels, drops, suppositories, sprays, liquids and powders. Conventional pharmaceutical carriers, aqueous, powder or oily bases, thickeners and the like may be necessary or desirable.

- 5 Compositions for oral administration include powders or granules, suspensions or solutions in water or non-aqueous media, capsules, sachets, or tablets. Thickeners, flavorings, diluents, emulsifiers, dispersing aids or binders may be desirable.

- Some of the compositions may potentially be administered as a pharmaceutically acceptable acid- or base- addition salt, formed by reaction with inorganic acids such as
- 10 hydrochloric acid, hydrobromic acid, perchloric acid, nitric acid, thiocyanic acid, sulfuric acid, and phosphoric acid, and organic acids such as formic acid, acetic acid, propionic acid, glycolic acid, lactic acid, pyruvic acid, oxalic acid, malonic acid, succinic acid, maleic acid, and fumaric acid, or by reaction with an inorganic base such as sodium hydroxide, ammonium hydroxide, potassium hydroxide, and organic bases such as mono-, di-, trialkyl and aryl amines and substituted
- 15 ethanolamines.

(2) Therapeutic Uses

- The dosage ranges for the administration of the compositions are those large enough to produce the desired effect in which the symptoms disorder are effected. The dosage should not be so large as to cause adverse side effects, such as unwanted cross-reactions, anaphylactic reactions,
- 20 and the like. Generally, the dosage will vary with the age, condition, sex and extent of the disease in the patient and can be determined by one of skill in the art. The dosage can be adjusted by the individual physician in the event of any counterindications. Dosage can vary, and can be administered in one or more dose administrations daily, for one or several days.

- Other compositions which do not have a specific pharmaceutical function, but which may
- 25 be used for tracking changes within cellular chromosomes or for the delivery of diagnostic tools for example can be delivered in ways similar to those described for the pharmaceutical products.

i) Chips and micro arrays

- Disclosed are chips where at least one address is the sequences or part of the sequences set forth in any of the nucleic acid sequences disclosed herein. Also disclosed are chips where at least
- 30 one address is the sequences or portion of sequences set forth in any of the peptide sequences disclosed herein.

 Also disclosed are chips where at least one address is a variant of the sequences or part of the sequences set forth in any of the nucleic acid sequences disclosed herein. Also disclosed are chips where at least one address is a variant of the sequences or portion of sequences set forth in

any of the peptide sequences disclosed herein.

k) Computer readable mediums

It is understood that the disclosed nucleic acids and proteins can be represented as a sequence consisting of the nucleotides of amino acids. There are a variety of ways to display these sequences, for example the nucleotide guanosine can be represented by G or g. Likewise the amino acid valine can be represented by Val or V. Those of skill in the art understand how to display and express any nucleic acid or protein sequence in any of the variety of ways that exist, each of which is considered herein disclosed. Specifically contemplated herein is the display of these sequences on computer readable mediums, such as, commercially available floppy disks, tapes, chips, hard drives, compact disks, and video disks, or other computer readable mediums. Also disclosed are the binary code representations of the disclosed sequences. Those of skill in the art understand what computer readable mediums. Thus, computer readable mediums on which the nucleic acids or protein sequences are recorded, stored, or saved.

Disclosed are computer readable mediums comprising the sequences and information regarding the sequences set forth herein. Also disclosed are computer readable mediums comprising the sequences and information regarding the sequences set forth herein wherein the sequences do not include SEQ ID NOs: SEQ ID NOs:1-19.

l) Kits

Disclosed herein are kits that are drawn to reagents that can be used in practicing the methods disclosed herein. The kits can include any reagent or combination of reagent discussed herein or that would be understood to be required or beneficial in the practice of the disclosed methods. For example, the kits could include primers to perform the amplification reactions discussed in certain embodiments of the methods, as well as the buffers and enzymes required to use the primers as intended. For example, disclosed is a kit for assessing a subject's risk for acquiring cancer, such as colon cancer, comprising the primers or probes that hybridize to the sequences set forth in SEQ ID NOs: 1, 3, 5, 7, 9, 11, and 13, for example.

m) Compositions with similar functions

It is understood that the compositions disclosed herein have certain functions, such as inhibiting gamma2 function or binding alpha6 integrin or inhibiting beta4 function. Disclosed herein are certain structural requirements for performing the disclosed functions, and it is understood that there are a variety of structures which can perform the same function which are related to the disclosed structures, and that these structures will ultimately achieve the same result, for example stimulation or inhibition alpha6beta4 signaling or interruption of the alpha6beta4 signaling pathway.

2. Methods of making the compositions

The compositions disclosed herein and the compositions necessary to perform the disclosed methods can be made using any method known to those of skill in the art for that particular reagent or compound unless otherwise specifically noted.

5 a) Nucleic acid synthesis

For example, the nucleic acids, such as, the oligonucleotides to be used as primers can be made using standard chemical synthesis methods or can be produced using enzymatic methods or any other known method. Such methods can range from standard enzymatic digestion followed by nucleotide fragment isolation (see for example, Sambrook *et al.*, *Molecular Cloning: A Laboratory*
10 *Manual*, 2nd Edition (Cold Spring Harbor Laboratory Press, Cold Spring Harbor, N.Y., 1989) Chapters 5, 6) to purely synthetic methods, for example, by the cyanoethyl phosphoramidite method using a Milligen or Beckman System 1Plus DNA synthesizer (for example, Model 8700 automated synthesizer of Milligen-Bioscience, Burlington, MA or ABI Model 380B). Synthetic methods useful for making oligonucleotides are also described by Ikuta *et al.*, *Ann. Rev. Biochem.*
15 53:323-356 (1984), (phosphotriester and phosphite-triester methods), and Narang *et al.*, *Methods Enzymol.*, 65:610-620 (1980), (phosphotriester method). Protein nucleic acid molecules can be made using known methods such as those described by Nielsen *et al.*, *Bioconjug. Chem.* 5:3-7 (1994).

b) Peptide synthesis

20 One method of producing the disclosed proteins, or fragments of the disclosed proteins, such as a fragment of SEQ ID NO:6, is to link two or more peptides or polypeptides together by protein chemistry techniques. For example, peptides or polypeptides can be chemically synthesized using currently available laboratory equipment using either Fmoc (9-fluorenylmethyloxycarbonyl) or Boc (*tert*-butyloxycarbonyl) chemistry. (Applied Biosystems,
25 Inc., Foster City, CA). One skilled in the art can readily appreciate that a peptide or polypeptide corresponding to the disclosed proteins, for example, can be synthesized by standard chemical reactions. For example, a peptide or polypeptide can be synthesized and not cleaved from its synthesis resin whereas the other fragment of a peptide or protein can be synthesized and subsequently cleaved from the resin, thereby exposing a terminal group which is functionally
30 blocked on the other fragment. By peptide condensation reactions, these two fragments can be covalently joined via a peptide bond at their carboxyl and amino termini, respectively, to form an antibody, or fragment thereof. (Grant GA (1992) *Synthetic Peptides: A User Guide*. W.H. Freeman and Co., N.Y. (1992); Bodansky M and Trost B., Ed. (1993) *Principles of Peptide Synthesis*. Springer-Verlag Inc., NY (which is herein incorporated by reference at least for
35 material related to peptide synthesis). Alternatively, the peptide or polypeptide is independently synthesized *in vivo* as described herein. Once isolated, these independent peptides or polypeptides

may be linked to form a peptide or fragment thereof via similar peptide condensation reactions.

For example, enzymatic ligation of cloned or synthetic peptide segments allow relatively short peptide fragments to be joined to produce larger peptide fragments, polypeptides or whole protein domains (Abrahmsen L et al., Biochemistry, 30:4151 (1991)). Alternatively, native
5 chemical ligation of synthetic peptides can be utilized to synthetically construct large peptides or polypeptides from shorter peptide fragments. This method consists of a two step chemical reaction (Dawson et al. Synthesis of Proteins by Native Chemical Ligation. Science, 266:776-779 (1994)).

The first step is the chemoselective reaction of an unprotected synthetic peptide--thioester with another unprotected peptide segment containing an amino-terminal Cys residue to give a thioester-
10 linked intermediate as the initial covalent product. Without a change in the reaction conditions, this intermediate undergoes spontaneous, rapid intramolecular reaction to form a native peptide bond at the ligation site (Baggiolini M et al. (1992) FEBS Lett. 307:97-101; Clark-Lewis I et al., J.Biol.Chem., 269:16075 (1994); Clark-Lewis I et al., Biochemistry, 30:3128 (1991); Rajarathnam K et al., Biochemistry 33:6623-30 (1994)).

15 Alternatively, unprotected peptide segments are chemically linked where the bond formed between the peptide segments as a result of the chemical ligation is an unnatural (non-peptide) bond (Schnolzer, M et al. Science, 256:221 (1992)). This technique has been used to synthesize analogs of protein domains as well as large amounts of relatively pure proteins with full biological activity (deLisle Milton RC et al., Techniques in Protein Chemistry IV. Academic Press, New
20 York, pp. 257-267 (1992)).

The disclosed proteins and polypeptides such as that for SEQ ID NO:6, beta4 integrin, can be made using any traditional recombinant biotechnology method. Examples of such methods can be found in Sambrook et al. which is herein incorporated by reference at least for material related to production of proteins and antibodies.

25 c) Processes for making the compositions

Disclosed are processes for making the compositions as well as making the intermediates leading to the compositions. For example, disclosed are nucleic acids in SEQ ID NOs:1, 3, 5, 7, 9, 11, and 13. There are a variety of methods that can be used for making these compositions, such as synthetic chemical methods and standard molecular biology methods. It is understood that the
30 methods of making these and the other disclosed compositions are specifically disclosed.

Disclosed are nucleic acid molecules produced by the process comprising linking in an operative way a nucleic acid molecule comprising the sequence set forth in SEQ ID Nos:1, 3, 5, 7, 9, 11, or 13 or a fragment thereof, and a sequence controlling the expression of the nucleic acid.

Also disclosed are nucleic acid molecules produced by the process comprising linking in

an operative way a nucleic acid molecule comprising a sequence having at least 80% identity to a sequence set forth in SEQ ID Nos:1, 3, 5, 7, 9, 11, or 13 or a fragment thereof, and a sequence controlling the expression of the nucleic acid.

Disclosed are nucleic acid molecules produced by the process comprising linking in an
5 operative way a nucleic acid molecule comprising a sequence that hybridizes under stringent hybridization conditions to a nucleic acid molecule comprising the sequence set forth in SEQ ID Nos:1, 3, 5, 7, 9, 11, or 13 or a fragment thereof, and a sequence controlling the expression of the nucleic acid.

Disclosed are nucleic acid molecules produced by the process comprising linking in an
10 operative way a nucleic acid molecule comprising a sequence encoding a peptide set forth in SEQ ID Nos:2, 4, 6, 8, 10, 12, or 14 or a fragment thereof, and a sequence controlling an expression of the nucleic acid molecule.

Disclosed are nucleic acid molecules produced by the process comprising linking in an
operative way a nucleic acid molecule comprising a sequence encoding a peptide having 80%
15 identity to a peptide set forth in SEQ ID Nos:2, 4, 6, 8, 10, 12, or 14 or a fragment thereof and a sequence controlling an expression of the nucleic acid molecule.

Disclosed are nucleic acids produced by the process comprising linking in an operative
way a nucleic acid molecule comprising a sequence encoding a peptide having 80% identity to a
peptide set forth in SEQ ID Nos:2, 4, 6, 8, 10, 12, or 14 or a fragment thereof, wherein any change
20 from the sequences set forth in SEQ ID Nos:2, 4, 6, 8, 10, 12, or 14 or a fragment thereof are conservative changes and a sequence controlling an expression of the nucleic acid molecule.

Disclosed are cells produced by the process of transforming the cell with any of the
disclosed nucleic acids. Disclosed are cells produced by the process of transforming the cell with
any of the non-naturally occurring disclosed nucleic acids.

25 Disclosed are any of the disclosed peptides produced by the process of expressing any of the disclosed nucleic acids. Disclosed are any of the non-naturally occurring disclosed peptides produced by the process of expressing any of the disclosed nucleic acids. Disclosed are any of the disclosed peptides produced by the process of expressing any of the non-naturally disclosed nucleic acids.

30 Disclosed are animals produced by the process of transfecting a cell within the animal with any of the nucleic acid molecules disclosed herein. Disclosed are animals produced by the process of transfecting a cell within the animal any of the nucleic acid molecules disclosed herein, wherein the animal is a mammal. Also disclosed are animals produced by the process of transfecting a cell within the animal any of the nucleic acid molecules disclosed herein, wherein the mammal is

mouse, rat, rabbit, cow, sheep, pig, or primate.

Also disclosed are animals produced by the process of adding to the animal any of the cells disclosed herein.

d) Products produced from selection protocols

5 Also disclosed are methods of obtaining molecules that act as functional regulators of integrin function, integrin receptor function, and functional regulators of signaling pathways related to integrin receptors, in particular integrin alpha6beta4.

Disclosed are methods for isolating molecules that interact with the proteins set forth in SEQ ID Nos:2, 4, 6, 8, 10, 12, or 14 or a fragment thereof comprising, interacting a library of
10 molecules with the proteins set forth in SEQ ID Nos:2, 4, 6, 8, 10, 12, or 14 or a fragment thereof, removing the unbound molecules, and collecting the molecules that are bound to at least one of the proteins set forth in SEQ ID Nos:2, 4, 6, 8, 10, 12, or 14 or a fragment thereof.

3. Methods of using the compositions

a) Methods of using the compositions as research tools

15 The compositions can be used for example as targets in combinatorial chemistry protocols or other screening protocols to isolate molecules that possess desired functional properties related to A6B4 and A6B1 signaling pathways.

The disclosed compositions can also be used diagnostic tools related to diseases such as cancer, such as colon cancer.

20 The disclosed compositions can be used as discussed herein as either reagents in micro arrays or as reagents to probe or analyze existing microarrays. The disclosed compositions can be used in any known method for isolating or identifying single nucleotide polymorphisms. The compositions can also be used in any method for determining allelic analysis of for example, beta4 alleles having varying function, particularly allelic analysis as it relates to beta4 signaling and
25 functions. The compositions can also be used in any known method of screening assays, related to chip/micro arrays. The compositions can also be used in any known way of using the computer readable embodiments of the disclosed compositions, for example, to study relatedness or to perform molecular modeling analysis related to the disclosed compositions.

b) Methods for affecting cancer

30 The disclosed compositions can be used to affect the growth of cancer cells because as disclosed herein, the disclosed relationships are fundamental to the ability of cancer cells to continue growing. The disclosed compositions, such as antisense constructs that will inhibit the production of either alpha6, beta4, or the gamma2 chain of laminin5 reduce the proliferation of cancer cells. In fact, the compositions cannot only reduce the proliferation of the cancer cells, but

the compositions can kill the cancer cells, as shown herein.

It is understood that cancer is caused by a variety of cellular events, of which certain events related to alpha6 integrin (up regulated), beta4 integrin (upregulated) and gamma2 (upregulated) allow the continued viability of cancer cells, and that interference of these upregulated molecules inhibits the growth and kills the cancer cells. However, there are other known events that can cause non-cancerous cells to become oncogenic. For example, Abl, Ras, EGF receptor, Erb-2, APC, beta-catenin, Arf, Mdm2, p53, Rb, Myc are known to be involved in oncogenesis, and some of these molecules (For example, Ras, APC loss, p53 loss) are directly related to the disclosed target signal transduction pathways mediated by the alpha6beta4 receptor. Just as the presently disclosed compositions can be used as therapeutics targeting the disclosed relationships, so there are other targets (For example, Abl, Erb-2) for which pharmaceutical compositions have been developed (For example, Glivec, Herceptin, respectively).

It is understood that the disclosed anti-cancer compositions can be used in combination with other anti-cancer compositions. Great benefits can be obtained from using anti-cancer compositions that target different molecules, in the same signal transduction pathway as well as, or in addition to, targeting molecules in different signal transduction pathways than those disclosed herein. Thus, the disclosed compositions which can affect the growth of cancer cells, indeed kill cancer cells, can be used in conjunction with any other chemotherapy, radiation, or any other anti-cancer therapy.

Disclosed are methods of reducing the proliferation of a cancer cell which comprises inhibiting ligand binding to an integrin receptor on the cancer cell, wherein the integrin receptor comprises an integrin.

Also disclosed are methods of reducing the proliferation of a cancer cell which comprises reducing integrin-integrin interaction, integrin receptor clustering interaction, or integrin-non-integrin protein interaction.

Also disclosed are methods of selectively reducing the proliferation of cancer cells which comprises reducing integrins from interacting with one another, integrins from clustering, or integrins from interacting with other proteins associated with cancer cells.

Disclosed are methods of reducing the proliferation of a cancer cell which comprises reducing the production of an integrin by the cancer cell.

Also disclosed are methods of reducing the proliferation of a cancer cell which comprises reducing the production of an integrin receptor ligand by the cancer cell.

Disclosed are methods of reducing the proliferation of a cancer cell which comprises

interfering with an integrin signaling pathway.

Also disclosed are methods of selectively killing or reducing the proliferation of cancer cells which comprises inhibiting ligand binding to integrin receptors on the cancer cells, wherein the integrin receptor comprises a B4 integrin.

5 Disclosed are methods of selectively killing or reducing the proliferation of cancer cells which comprises inhibiting ligand binding to integrin receptors on the cancer cells, wherein the ligand comprises a laminin5.

Also disclosed are methods of selectively killing or reducing the proliferation of cancer cells which comprises inhibiting ligand binding to integrin receptors on the cancer cells, wherein
10 the ligand comprises the gamma2 subunit of the laminin L5.

Disclosed are methods of selectively killing or reducing the proliferation of cancer cells which comprises inhibiting ligand binding to integrin receptors on the cancer cells, wherein the integrin receptor comprises an alpha6 integrin.

Further disclosed are methods of selectively killing or reducing the proliferation of cancer
15 cells which comprises preventing integrin receptor subunits from interacting with one another, preventing integrin clustering, or preventing integrin receptor subunits from interacting with other proteins on cancer cells, wherein the integrin receptor comprises a B4 integrin.

Disclosed are methods of selectively killing or reducing the proliferation of cancer cells which comprises reducing the production of laminin by the cancer cells, wherein the laminin
20 comprises laminin5, and/or any of the subunits of laminin5, such as gamma2.

Also disclosed is a method of selectively killing or reducing the proliferation of cancer cells which comprises interfering with an integrin signaling pathway, wherein the integrin signaling pathway comprises a B4 integrin or an alpha6 integrin or a beta1 integrin or a laminin or a laminin5 or the gamma2 subunit of laminin5.

25 Also disclosed are methods, wherein the integrin receptor comprises integrin B4 and/or wherein the integrin receptor comprises integrin A6, and/or wherein the ligand that binds to the integrin receptor is laminin5, and/or wherein the integrin receptor is A6B4, and/or wherein the ligand comprises laminin5, and/or wherein the ligand comprises the gamma-2 subunit of laminin5, and/or wherein inhibiting ligand binding to an integrin receptor does not occur by using an
30 antisense molecule to A6, and/or wherein inhibiting ligand binding to an integrin receptor comprises contacting a A6 integrin with a composition that inhibits ligand binding, and/or wherein inhibiting ligand binding to an integrin receptor comprises contacting a B4 integrin with a composition that inhibits ligand binding, and/or wherein inhibiting ligand binding to an integrin receptor comprises contacting a laminin5 with a composition that inhibits ligand binding, and/or

wherein inhibiting ligand binding to an integrin receptor comprises contacting a gamma-2 subunit
 with a composition that inhibits ligand binding, and/or wherein reducing integrin-integrin
 interaction, integrin receptor clustering interaction, or integrin-non-integrin protein interaction
 does not occur by using a B4-delta-cyt, and/or wherein reducing integrin-integrin interaction,
 5 integrin receptor clustering interaction, or integrin-non-integrin protein interaction does not occur
 by using an antisense molecule to A6, and/or wherein reducing integrin-integrin interaction,
 integrin receptor clustering interaction, or integrin-non-integrin protein interaction comprises
 contacting a A6 integrin with a composition that inhibits an interaction between the B4 integrin
 and another integrin or protein molecule, and/or wherein reducing integrin-integrin interaction,
 10 integrin receptor clustering interaction, or integrin-non-integrin protein interaction comprises
 contacting a B4 integrin with a composition that inhibits the interaction between the B4 integrin
 and another integrin or protein molecule, and/or wherein reducing integrin-integrin interaction,
 integrin receptor clustering interaction, or integrin-non-integrin protein interaction comprises
 contacting a laminin5 with a composition that inhibits ligand binding, and/or wherein reducing
 15 integrin-integrin interaction, integrin receptor clustering interaction, or integrin-non-integrin
 protein interaction comprises contacting a gamma2 subunit with a composition that inhibits ligand
 binding, and/or wherein the non-integrin protein comprises a growth factor receptor, and/or
 wherein the non-integrin protein comprises a hemi-desmosome junction, and/or wherein the non-
 integrin protein comprises a SH2 domain, and/or wherein the non-integrin protein comprises a Shc
 20 protein, and/or wherein the non-integrin protein comprises a IRS-1 protein, and/or wherein the
 non-integrin protein comprises a IRS-2 protein, and/or wherein reducing the production of an
 integrin does not occur by using an antisense molecule to A6, and/or wherein the production of an
 integrin is reduced by inhibiting signaling leading to induction of expression of an integrin, and/or
 wherein reducing the production of an integrin comprises inhibiting alpha6 production, and/or
 25 wherein inhibiting alpha6 production further comprises using antisense molecules to alpha6
 mRNA, and/or wherein reducing the production of an integrin comprises inhibiting beta4
 production, and/or wherein inhibiting beta4 production further comprises using antisense
 molecules to beta4 mRNA, and/or wherein reducing the production of an integrin receptor ligand
 comprises inhibiting gamma2 production, and/or wherein reducing the production of an integrin
 30 receptor ligand comprises inhibiting laminin production, and/or wherein reducing the production
 of an integrin receptor ligand comprises inhibiting laminin5 production, and/or wherein interfering
 with an integrin signaling pathway does not occur by using a B4-delta-cyt, and/or wherein
 interfering with an integrin signaling pathway does not occur by using an antisense molecule to
 A6, and/or wherein interfering with an integrin signaling pathway comprises contacting an A6
 35 integrin in the cell with a composition that inhibits ligand binding, and/or wherein interfering with
 an integrin signaling pathway comprises contacting a B4 integrin with a composition that inhibits

ligand binding, and/or wherein interfering with an integrin signaling pathway comprises contacting a laminin5 with a composition that inhibits ligand binding, and/or wherein interfering with an integrin signaling pathway comprises contacting a gamma2 subunit with a composition that inhibits ligand binding, and/or wherein interfering with an integrin signaling pathway comprises
 5 contacting the cancer cell with a molecule that interferes with at least one of talin, paxillin, vinculin, a CAS family protein, CRX, NCK, FAK, ILK, Src, Fyn, Shc, Grb-2, Guanine nucleotide exchange factors, SOS, DOCK 180, Vav, Syk, P-1-3 kinase, AKT, Bad, Bid, Caspase 9, Cdc42, PAK, Rac, Rho, Rho kinase, Ras, Caveolin, Tetraspan, Receptor-type protein tyrosine phosphatase, SHP-2, Alpha-actinin, Filamin, Cytohesin, Beta3-endonexin, ICAP-1, RACK-1, CIB,
 10 actin, receptor tyrosine kinase, IRS-1 or IRS-2, and/or wherein interfering with an integrin signaling pathway comprises contacting the cancer cell with an agent that interferes with post-translational modification of integrins, and/or wherein the post translational modification is glycosylation or phosphorylation.

Also disclosed are methods, wherein the integrin comprises an A6 integrin, and/or wherein
 15 the integrin comprises a B4 integrin, and/or further comprising reducing a laminin5-integrin interaction, and/or further comprising reducing a laminin5 gamma2 integrin interaction, and/or wherein the cancer cell comprises normal p53, and/or wherein the proliferation of the cancer cells is not dependent on AKT/PKB, and/or wherein reducing the proliferation of the cancer cells is selective, and/or wherein the cancer cell is not an MDA-MB-435 cell, and/or wherein the cancer
 20 cell is not an HMT-3522 cell, and/or wherein the cancer cell is not an RKO colon carcinoma line, and/or wherein the cancer cell does not express exogenous B4 integrin, and/or further comprising contacting the cancer cells with a small molecule, peptide, peptide mimetic, or oligonucleotide or synthetic analog thereof, and/or wherein the cancer cells are contacted with dominant-negative beta 4 integrin, and/or wherein the cancer cell is contacted with an antisense molecule, and/or
 25 wherein the antisense molecule is linked to a leader sequence which enables translocation across a cell membrane, and/or wherein the leader sequence binds to a cell surface protein which facilitates internalization, and/or wherein the leader sequence is TAT or antennapedia, or fragment thereof, and/or wherein the antisense molecule is an alpha6 RNA antisense molecule, and/or wherein the small molecule peptide, peptide mimetic, or oligonucleotide or synthetic analog thereof is linked
 30 to a carrier, and/or wherein the carrier is at least one of a lipidic carrier, charged carrier, retroviral carrier, TAT or fragment thereof, antennapedia or fragment thereof, or polyethylene glycol, and/or further comprising contacting the cancer cell with another agent which modulates cell signaling, a chemotherapeutic drug, or treated with radiation or angiogenesis inhibitor, and/or wherein reducing the proliferation of cancer cell is *in vitro*, and/or wherein reducing the proliferation of the
 35 cancer cell is *in vivo*, and/or wherein the cancer cell is selected from the group consisting of melanoma, adenoma, lymphoma, myeloma, carcinoma, plasmocytoma, sarcoma, glioma, thymoma,

leukemia, skin cancer, retinal cancer, breast cancer, prostate cancer, colon cancer, esophageal cancer, stomach cancer, pancreas cancer, brain tumors, lung cancer, ovarian cancer, cervical cancer, hepatic cancer, gastrointestinal cancer, and head and neck cancer cells, and/or wherein the cancer cell is killed, and/or wherein the cancer cell expresses a mutated Ras, and/or wherein the cancer
5 cell expresses a mutated Ras and a mutated p53, and/or wherein the cancer cell expresses a mutated Ras and activates the AKT/PKB protein, and/or wherein the cancer cell expresses a mutated Ras, a mutated p53, and activates the AKT/PKB protein, and/or wherein the cancer cell expresses a mutated APC, and/or wherein the cancer cell expresses a mutated Ras and mutated APC.

10 Disclosed are methods of reducing the proliferation of a cancer cell in a patient which comprises administering to the patient a composition which inhibits ligand binding to an integrin receptor on the cancer cell, wherein the integrin receptor comprises an integrin.

Also disclosed are methods of reducing the proliferation of a cancer cell in a patient which comprises administering to the patient a composition which reduces integrin-integrin interaction,
15 integrin receptor clustering interaction, or integrin-non-integrin protein interaction.

Also disclosed are methods of reducing the proliferation of a cancer cell in a patient which comprises administering to the patient a composition which reduces the production of an integrin or laminin by the cancer cell.

Further disclosed are methods of reducing the proliferation of a cancer cell in a patient
20 which comprises administering to the patient a composition which interferes with an integrin signaling pathway.

Also disclosed are methods, wherein the reduction in cancer cell proliferation is selective, and/or wherein the administering is local or systemic, and/or wherein the patient is additionally administered an agent which modulates cell signaling, a chemotherapeutic drug, or treated with
25 radiation or angiogenesis inhibitor, and/or wherein the additional agent is administered serially or in combination, and/or wherein the local administering is direct application to cancer cells, and/or wherein the direct application to cancer cells is performed during surgery, and/or wherein the direct application to cancer cells is performed topically to cancerous tissue, and/or wherein the systemic administration is by subcutaneous, intraperitoneal, intra-arterial, intravenous, or bolus
30 administration, or by application through a catheter or similar apparatus, and/or wherein the systemic administration comprises a long-term release formulation, and/or wherein the systemic administration is by oral administration, and/or wherein the oral administration comprises administering a pill, capsule, tablet, liquid or suspension.

Disclosed are methods of reducing the proliferation of a cancer cell in a patient which

comprises administering to a patient a vector comprising coding sequence for a protein or peptide which inhibits ligand binding to integrin receptors on cancer cells wherein the coding sequence is under the control of a promoter which functions in mammalian cells.

Also disclosed are methods of reducing the proliferation of a cancer cell in a patient which
5 comprises administering to a patient a vector comprising coding sequence for a protein or peptide which prevents integrin receptor subunits from interacting with one another, prevents integrin receptor clustering interaction, or prevents integrin receptor subunits from interacting with other proteins on cancer cells wherein the coding sequence is under the control of a promoter which functions in mammalian cells.

10 Further disclosed are methods of reducing the proliferation of a cancer cell in a patient which comprises administering to a patient a vector comprising coding sequence for a protein or peptide which prevents integrin receptor subunits from interacting with one another, prevents integrin receptor clustering interaction, or prevents integrin receptor subunits from interacting with other proteins in cancer cells wherein the coding sequence is under the control of a promoter which
15 functions in mammalian cells.

Also disclosed are method of reducing the proliferation of a cancer cell in a patient which comprises administering to a patient a vector comprising coding sequence for a protein or peptide which interferes with integrin subunit or laminin production wherein the coding sequence is under the control of a promoter which functions in mammalian cells.

20 Further disclosed are methods of reducing the proliferation of a cancer cell in a patient which comprises administering to a patient a vector comprising coding sequence for a protein or peptide which interferes with an integrin signaling pathway of cells wherein the coding sequence is under the control of a promoter which functions in mammalian cells.

Also disclosed are methods, wherein the reduction in cancer cell proliferation is selective,
25 and/or wherein the vector is administered directly to a cancer cell, and/or wherein the vector is administered directly to a normal cell, and/or wherein the vector is packaged in a viral vector or liposome, and/or wherein the vector is a retroviral vector, and/or wherein the vector is administered systemically, and/or wherein the direct administration is by topical application, and/or wherein the direct administration is by topical application, and/or wherein the direct
30 administration is performed during surgery, and/or wherein the direct administration is performed during surgery, and/or wherein the patient is an animal, such as a mammal, mouse, rabbit, primate, chimp, ape, gorilla, and human, and/or wherein the cancer cells are selected from the group consisting of melanoma, adenoma, lymphoma, myeloma, carcinoma, plasmocytoma, sarcoma, glioma, thyoma, leukemia, skin cancer, retinal cancer, breast cancer, prostate cancer, colon cancer,
35 esophageal cancer, stomach cancer, pancreas cancer, brain tumors, lung cancer, ovarian cancer,

cervical cancer, hepatic cancer, gastrointestinal cancer, and head and neck cancer cells, and/or wherein the patient is additionally administered at least one of another agent which modulates cell signaling, a chemotherapeutic drug, an angiogenesis inhibitor or treated with radiation, and/or wherein the other agent which modifies cell signaling, chemotherapeutic drug, angiogenesis
5 inhibitor or radiation treatment is administered serially or in combination.

c) Methods of gene modification and gene disruption

The disclosed compositions and methods can be used in targeted gene disruption and modification in any animal that can undergo these events. Gene modification and gene disruption refer to the methods, techniques, and compositions that surround the selective removal or
10 alteration of a gene or stretch of chromosome in an animal, such as a mammal, in a way that propagates the modification through the germ line of the mammal. In general, a cell is transformed with a vector which is designed to homologously recombine with a region of a particular chromosome contained within the cell, as for example, described herein. This homologous recombination event can produce a chromosome which has exogenous DNA
15 introduced, for example in frame, with the surrounding DNA. This type of protocol allows for very specific mutations, such as point mutations, to be introduced into the genome contained within the cell. Methods for performing this type of homologous recombination are disclosed herein.

One of the preferred characteristics of performing homologous recombination in
20 mammalian cells is that the cells should be able to be cultured, because the desired recombination event occur at a low frequency.

Once the cell is produced through the methods described herein, an animal can be produced from this cell through either stem cell technology or cloning technology. For example, if the cell into which the nucleic acid was transfected was a stem cell for the organism, then this cell,
25 after transfection and culturing, can be used to produce an organism which will contain the gene modification or disruption in germ line cells, which can then in turn be used to produce another animal that possesses the gene modification or disruption in all of its cells. In other methods for production of an animal containing the gene modification or disruption in all of its cells, cloning technologies can be used. These technologies generally take the nucleus of the transfected cell and
30 either through fusion or replacement fuse the transfected nucleus with an oocyte which can then be manipulated to produce an animal. The advantage of procedures that use cloning instead of ES technology is that cells other than ES cells can be transfected. For example, a fibroblast cell, which is very easy to culture can be used as the cell which is transfected and has a gene modification or disruption event take place, and then cells derived from this cell can be used to
35 clone a whole animal. Conditional knockouts can also be made which will conditionally delete

expression of the desired molecule, for example, an A6 or integrin or the gamma2 chain of laminin5.

d) Methods of diagnosing cancer

Methods of diagnosing cancer using the disclosed information and the disclosed
5 molecules. In particular disclosed are methods of diagnoses that rely on the combined upregulation of both the ligand and the cognate integrin receptor or cognate integrins. It is understood that all of the methods of diagnosis disclosed herein, can be used with any of the disclosed compositions, but they also can be used in conjunction, where for example, both the ligand and the integrin receptor are monitored and correlated with cancer developing.

10 Disclosed are methods of assessing a subject's risk of developing cancer comprising determining the amount of A6 present in a target cell obtained from the subject, wherein a determination of increased levels of A6 correlates with an increased risk of cancer.

Disclosed are methods of assessing a subject's risk of acquiring cancer comprising determining the amount of B4 present in a target cell obtained from the subject, wherein a
15 determination of increased levels of B4 correlates with an increased risk of cancer.

Disclosed are methods of assessing a subject's risk of acquiring cancer comprising determining the amount of laminin5 present in a target cell obtained from the subject, wherein a determination of increased levels of laminin5 correlates with an increased risk of cancer.

Disclosed are methods of assessing a subject's risk of acquiring cancer comprising
20 determining the amount of gamma2 subunit present in a target cell obtained from the subject, wherein a determination of increased levels of gamma2 subunit correlates with an increased risk of cancer.

Also disclosed are methods, further comprising comparing the amount A6 present to the amount in a control cell, and wherein determining the amount of A6 present in the target cell
25 comprises assaying the amount of A6 mRNA in the cell, and/or wherein the assaying the amount of mRNA in the cell comprises hybridizing a A6 probe to a sample of the subject's mRNA, and/or wherein the assaying the amount of mRNA in the cell comprises hybridizing a A6 primer to a sample of the subject's mRNA, and/or wherein the assaying the amount of mRNA further comprises performing an nucleic acid amplification reaction involving the primer, and/or wherein
30 the nucleic acid amplification reaction comprises reverse transcription, producing a cDNA, and/or wherein the nucleic acid amplification reaction further comprises performing a polymerase chain reaction on the cDNA, and/or wherein determining the amount of A6 present in the target cell comprises assaying the amount of A6 protein in the cell, and/or further comprising comparing the amount B4 present to the amount in a control cell, and/or further comprising comparing the

amount laminin5 present to the amount in a control cell, and/or further comprising comparing the amount gamma2 subunit present to the amount in a control cell.

C. Examples

Throughout this application, various publications are referenced. The disclosures of these
5 publications in their entireties are hereby incorporated by reference into this application in order to more fully describe the state of the art to which this invention pertains. The references disclosed are also individually and specifically incorporated by reference herein for the material contained in them that is discussed in the sentence in which the reference is relied upon.

It will be apparent to those skilled in the art that various modifications and variations can
10 be made in the present invention without departing from the scope or spirit of the invention. Other embodiments of the invention will be apparent to those skilled in the art from consideration of the specification and practice of the invention disclosed herein. It is intended that the specification and examples be considered as exemplary only, with a true scope and spirit of the invention being indicated by the following claims.

15 The following examples are put forth so as to provide those of ordinary skill in the art with a complete disclosure and description of how the compounds, compositions, articles, devices and/or methods claimed herein are made and evaluated, and are intended to be purely exemplary of the invention and are not intended to limit the scope of what the inventors regard as their invention. Efforts have been made to ensure accuracy with respect to numbers (e.g., amounts,
20 temperature, etc.), but some errors and deviations should be accounted for. Unless indicated otherwise, parts are parts by weight, temperature is in °C or is at ambient temperature, and pressure is at or near atmospheric.

1. Example 1 Colonic epithelial cells dependent on alpha6beta4 receptor signal transduction for growth in the absence of ECM

25 Integrin alpha6 and integrin beta4 and laminin gamma2 chains are essential for tumor cell survival in vivo. The data disclosed herein indicate that interfering with integrin alpha6-mediated signaling, integrin beta 4signaling, and laminin gamma2 signaling can constitute an effective approach to induce programmed cell death in cancer cells without damaging normal cells. Ablation of integrin alpha6-dependent signaling, integrin beta4-dependent signaling, and laminin
30 gamma2- dependent signaling in human cancer cell lines supports this.

The murine colonic epithelial transformation system introduced by D'Abaco et al. (1996) was used. In this system, control cells (control), cells containing an activated ras oncogene (Ras), a deletion in the APC gene (APCmin) or both alterations together (Ras+APCmin) can be compared with regard to their proliferation characteristics in tissue culture. The cells were derived from
35 transgenic mice containing a temperature-sensitive allele of SV40 large T under the control of a

gamma interferon-inducible promoter permitting conditional immortalization (Jat *et al.*, 1991). All experiments were carried out with the cells being kept at non-permissive temperature in the absence of gamma interferon. As shown previously (D'Abaco *et al.*, 1996), only the cells carrying activated Ras and the APCmin mutation were able to form colonies in soft agar in the absence of anchorage to a substratum. All other cell populations did not give rise to colonies under these conditions.

The frequency of tunel-staining cells (Fig.2A) and caspase 3 activity (Fig. 2B) in control, Ras, APCmin and Ras+APCmin colonic epithelial cells (D'Abaco *et al.*, 1996) in suspension was measured. Both assays show a strong suppression of apoptosis in Ras-APCmin cells, indicating that Ras and APCmin co-operate in preventing cell death in the absence of apparent ECM contacts. Activated Ras can protect cells from apoptosis via activation of the serine-threonine kinase AKT (Kauffmann-Zeh *et al.*, 1997; Khwaja *et al.*, 1997). However, both Ras and Ras+APCmin cells show equivalent levels of AKT phosphorylation (not shown), eliminating AKT as the relevant target of Ras+APCmin cooperation.

Although cancer cells have been thought to survive independently of integrin signaling, they frequently express high levels of alpha3 or alpha6 integrin receptors and/or their ligands (Dedhar *et al.*, 1993; Kennel *et al.*, 1989; Koshikawa *et al.*, 1999; Lohi *et al.*, 2000; Nejjarri *et al.*, 1999; Van Waes and Carey, 1992). These integrins are suspected to play a role in invasion and metastasis (Mukhopadhyay *et al.*, 1999; Shaw *et al.*, 1997). Ras and APCmin mutations would induce alterations in the mRNA expression patterns of integrin and ECM components. Induction of alpha6 integrin alone, was specific to Ras+APCmin cells (Fig. 3A). In addition, we found elevated expression of the laminin alpha5, gamma2 and beta2 chains induced in Ras and Ras+APCmin cells (Fig. 3B). These chains are components of laminins 5, 10 and 11 (Malinda and Kleinman, 1996), which are ligands of integrin receptors alpha6/beta4 and alpha3/beta1 (Kikkawa *et al.*, 2000; Niessen *et al.*, 1994). These results indicate that Ras and APCmin mutations can cooperate to induce autocrine activation of integrin receptors.

Alpha6 integrin can form functional receptors together with either beta1 or beta4 integrins (Clark and Brugge, 1995), leading to signaling through downstream effectors such as focal adhesion kinase (FAK) or the SH2 domain adapter Shc (Giancotti and Ruoslahti, 1999). Conversely, beta4 integrin only binds to alpha6 integrin (Clark and Brugge, 1995). In transformed colonic epithelial cells alpha6 integrin functions in conjunction with integrin beta4 and is engaged by the laminin gamma2 chain to activate Shc (see Fig. 4). Consistent with alpha6 integrin induction, only Ras+APCmin cells can bind a laminin gamma2-specific peptide in an alpha6 and beta4 integrin-specific manner. In contrast, an alpha3/beta1 and alpha6/beta1-specific peptide derived from the laminin gamma1 chain, only binds to control cells (Fig. 4A). In addition, only in

Ras+APCmin cells is Shc phosphorylated in response to clustering of the integrin alpha6/beta4 receptor by beta4-specific antibodies (Mainiero *et al.*, 1995) and by the laminin gamma2-specific peptide (Fig. 4B). In both cases p52^{Shc} is the major phosphorylated form. Moreover, alpha6 integrin is required for Shc activation, as Shc phosphorylation is inhibited in response to
5 expression of a dominant-negative beta4 integrin mutant. This mutant lacks the cytoplasmic signaling domain but selectively binds to alpha6 integrin to form a ligand binding, yet signaling-defective alpha6/beta4 integrin (Spinardi *et al.*, 1993) (Fig. 4B)

Anti-sense RNAs specific for alpha6 integrin and gamma2 laminin, as well as dominant-negative beta4 integrin, were expressed and results indicated that integrin alpha6, integrin beta4,
10 and laminin gamma2 expression are relevant for the survival of Ras+APCmin cells in Ras+APCmin, Ras, APC and control cells. All three constructs efficiently inhibit the growth of Ras+APCmin cells in soft agar (not shown) and when attached to plastic (Fig. 5A1). In contrast, the proliferation of wt, Ras and APCmin cells is not affected by this treatment (Fig. 5A1). Importantly, the colony formation of Ras+APC cells exposed to any of the three inhibitory
15 constructs can be rescued by co-expression of baculovirus p35, a potent inhibitor of caspase activity and of apoptosis (Resnicoff *et al.*, 1998) (Fig. 5A1). Similarly, co-expression of exogenous integrin alpha6 mRNA efficiently rescues Ras-APCmin cell proliferation inhibited by alpha6 anti-sense RNA (Fig. 5A1). This is mirrored by a rescue of alpha6 integrin expression on the cell surface of alpha6 anti-sense RNA expressing cells (Fig. 5B). Moreover, ectopic alpha6
20 integrin expression also rescues apoptosis induced by dominant-negative beta4 integrin (Fig. 5A1). The latter is expressed at constant levels (Fig. 5B). As one would expect, the cells expressing laminin gamma2 antisense RNA could not be rescued by integrin alpha6 over-expression, but showed significant rescue when plated on dishes coated with laminin gamma2 peptide (Fig. 5A1). Similarly, this peptide increases the survival of Ras+APCmin cells in
25 suspension, as indicated by lower caspase 3 activity (Fig. 2C). In summary, the survival of Ras-APCmin cells depends on the expression of alpha6/beta4 integrin and the laminin gamma2 chain. Conversely, the inhibition of these gene activities leads to selective killing of transformed cells.

Data also indicate that cell death due to lack of alpha6/beta4 integrin receptor signaling in Ras-APCmin cells is independent of the death signaling pathway involving the induction of
30 mitochondrial damage and caspase 9 activity. Instead, the data indicate an involvement of the Fas/TNF receptor/death domain protein/caspase8 pathway (Kruidering and Evan, 2000) in the control of tumor cell survival by alpha6 integrin-containing integrin receptors.

Control, APCmin and Ras cells, which lack alpha6/beta4 integrin receptor signaling activity, show high levels of caspase 8 activity when kept in suspension, while at the same time
35 caspase 9 activity cannot be detected (Fig. 2D).

Although Ras-APCmin cells can be rescued from apoptosis by caspase inhibitor baculovirus p35 (Fig. 5 A1, A2 and text above), cell death induced by expression of dominant-negative beta4 integrin cannot be prevented by expression of the survival factor Bcl2 (see Fig. 5 A2). Bcl2 binds to and neutralizes BH3-domain killer proteins that cause mitochondrial damage, cytochrome C release and caspase 9 activation (Luo *et al.*, 1998).

Ras cells which lack alpha6/beta4 integrin receptor signaling activity and cannot survive in the absence of ECM contacts, and Ras+APCmin cells which depend on alpha6/beta4 integrin receptor signaling for survival, show equivalent levels of AKT phosphorylation (data not shown). This indicates that AKT does not serve as the key target for alpha6/beta4 integrin receptor signaling. AKT has been described to promote cell survival via phosphorylation and inactivation of Bad, a BH3-domain killer protein (Datta *et al.*, 1999).

2. Example 2 Other cancer cells dependent on alpha6beta4 receptor signal transduction for growth in the absence of ECM

The survival of highly transformed primary mouse embryo fibroblasts expressing Ras and Myc oncoproteins in conjunction with a homozygous ARF null mutation (Kamijo *et al.*, 1997) depends on expression and function of alpha6 integrin (Fig. 5C), thus demonstrating that this principle also may apply to transformed mesenchymal cells. Moreover, the human colon cancer cell line SW480 carrying multiple oncogenic mutations such as activated Ras (Fujita *et al.*, 1988), amplified c-Myc (Cherif *et al.*, 1988), a mutated APC allele (Munemitsu *et al.*, 1995) and a p53 mutation (Abarzua *et al.*, 1995) is effectively killed by anti-sense integrin alpha6 and laminin gamma2 RNAs as well as dominant-negative beta4 integrin (Fig. 4D). Similarly, integrin alpha6 ablation even leads to apoptosis of Ras/APCmin cells in the presence of active SV40 large T (not shown). Thus the survival of different types of highly transformed cells depends on alpha6 integrin expression, beta4 integrin expression and laminin5 expression irrespective of the status of major tumor suppressor genes, such as arf, p53 or rb, and for at least sw480 cells, murine colon epithelial cells, and the ras/APCmin cells, alpha6, beta4, and laminin5/gamma2 are required. The sensitivity of transformed cells to ablation of integrin signaling, for example, A6B4 integrin receptor signaling, is thus quite remarkable in its apparent generality.

Oncogenic mutations co-operate to engage autocrine integrin signaling. Importantly, this signaling mechanism, involving alpha6 integrin, beta4 integrin and the laminin gamma2 chain, becomes an essential component of the survival mechanism in transformed colonic epithelial cells. In addition, fibroblasts and human colon cancer cells also rely on integrin alpha6 for survival. In contrast, normal and partially transformed cells that express alpha6 integrin at low levels do not require this polypeptide for survival. Thus, integrin signaling inhibition can lead to selective killing of cancer cells.

The laminin-integrin receptor signaling loop is also relevant to the survival of cells transformed by other combinations of oncogenic lesions. Introduction of activated Ras together with dominant-negative p53 (Lloyd et al., 1997) into murine colonic epithelial cells (D'Abaco *et al.*, 1996; see also above) via retroviral infection supports this. Four distinct polyclonal pools of
 5 infected cells were derived by drug selection: 1) Control (bleo/neo) cells, infected with two retroviruses carrying neomycin (neo) or bleomycin (bleo) drug resistance markers, respectively; 2) Ras cells, infected with a Ras/neo virus and a virus with the bleo marker; 3) DNp53 cells, infected with a DNp53/bleo virus and a virus with the neo marker; and 4) Ras/DNp53 cells, infected with a Ras/neo virus and a DNp53/bleo virus.

10 As the colon epithelial cells contain temperature-sensitive SV40 large T under control of the gamma interferon promoter, all cell populations were drug-selected at the permissive temperature (33°C) in the presence of gamma interferon. All further experiments were carried out at the non-permissive temperature for SV40 large T in the absence of gamma interferon, as described above for Ras+APCmin cells.

15 Similar to Ras+APCmin cells, only Ras/DNp53 cells but neither bleo/neo, Ras/bleo or DNp53/neo cells were able to grow in the absence of ECM contact in soft agar (not shown). Moreover, Ras and DNp53 cooperate in the suppression of caspase 3 activity in suspension (Fig. 2B), and in the induction of alpha6 integrin expression (Fig. 4A). In Ras/DNp53 cells passaged through a single round of soft agar growth, alpha6 integrin is expressed at even higher levels,
 20 demonstrating a correlation between the ability of the cells to survive in the absence of ECM contact and alpha6 integrin expression. In Ras/DNp53 cells beta4 integrin expression levels are also increased, when compared to controls (Fig. 4A). As expected expression of the laminin gamma2 chain is induced in Ras/bleo and Ras/DNp53 cells (Fig. 4B).

Anti-sense RNAs specific for alpha6 integrin and gamma2 laminin, as well as dominant-
 25 negative beta4 integrin were expressed in Ras/DNp53, Ras/bleo, DNp53/neo and neo/bleo cells. All three constructs efficiently inhibit the growth of Ras/DNp53 cells when attached to plastic. In contrast, the proliferation of neo/bleo, Ras/bleo and DNp53/neo cells is not affected by this treatment (Fig. 5A3). Anti-sense alpha6 integrin or dominant-negative beta4 integrin also inhibit the growth of Ras/DNp53 cells in soft agar (Fig. 5A4).

30 Integrin alpha6 expression can also be induced by activated Ras and DNp53 in primary murine colon crypt epithelial cells (not shown). Furthermore introduction of Ras and Myc oncoproteins into our control colon epithelial cells also leads to an induction of alpha6 expression (not shown), suggesting that induction of alpha6 integrin expression may be an integral component of distinct oncogene cooperation paradigms

a) Cell types to be tested

As described herein, there are a variety of cancer and transformed cell types that express alpha6 integrin and laminins at high levels (Dedhar *et al.*, 1993; Koshikawa *et al.*, 1999; Lohi *et al.*, 2000; Van Waes and Carey, 1992). As disclosed herein the following cell lines other than the
5 fibroblasts require alpha6 and beta4 integrin expression, as well as laminin gamma2 chain expression: (1) the human colon carcinoma cell line SW480 (data not shown), (2) Ras+APCmin-transformed murine colon epithelial cells (see Fig. 3A) Murine fibroblasts transformed by activated Ras and Myc in conjunction with homozygous Arf null mutation require alpha6 expression for their survival. These cells do not express integrin beta4, but express beta1 integrin,
10 indicating the importance of alpha6/beta1 receptors in the transformed fibroblasts.

b) Inducible expression of inhibitors

To interrupt alpha6 integrin and laminin gamma2 chain-dependent signaling in tumors regulatable alpha6 integrin and laminin gamma2 antisense mRNA expression as well as regulatable expression of the beta4 integrin the dominant-negative polypeptide described in
15 Section C were used. Constitutive expression of these inhibitors leads to rapid cell death in tissue culture. In contrast, the establishment of clonal cell lines with inducible expression of anti-sense mRNAs or dominant-negative beta4 integrin minimize such limitations.

The doxycycline (dox)-inducible reverse tetracycline transactivator (rtTA) is frequently used to overexpress transgenes in a temporally regulated fashion in vitro and in vivo (Efrat *et al.*,
20 1995; Gossen *et al.*, 1995; Ray *et al.*, 1997). These systems are, however, often compromised by the levels of gene expression in the absence of dox administration. The tetracycline controlled transcriptional silencer (tTS), a fusion protein containing the tet repressor and the KRAB-AB domain of the kid-1 transcriptional repressor, is inhibited by doxycycline. As shown in tissue culture (Freundlieb *et al.*, 1999) and in transgenic mice (Zhu *et al.*, 2001), tTS tightens the control
25 of transgene expression in rtTA-based systems, i.e. tTS effectively eliminates leaky baseline expression without altering the inducibility of rtTA-regulated genes. Thus optimal "off/on" regulation of gene expression can be accomplished with the combined use of tTS and rtTA. The complete expression system is commercially available from CLONTECH. This system can be used for the preparation of all inducible cell lines.

30 Another regulatable system comprising the estrogen-dependent transactivator GalER-VP16 and a promoter under the control of Gal4 DNA binding sites (Brasemann *et al.*, 1993) can be used. (Perez-Roger *et al.*, 1997). This particular experimental set up, was used herein to show that the induction of the beta4 integrin dominant-negative mutant in SW480 colon cancer cells induces apoptosis in vitro, as measured by caspase 3 activation (Fig. 2B). For in vivo use,
35 however, a point mutation has to be introduced into the GalER-VP16 transactivator that eliminates

its sensitivity to estrogen while retaining its response to the anti-estrogen 4OH-tamoxifen (Littlewood *et al.*, 1995). 4OH-tamoxifen has been demonstrated to reversibly regulate in vivo the activity of another regulatable transactivator, the MycER chimera (Pelengaris *et al.*, 1999).

Typically, inducible cell lines are first stably transfected with the anti-sense or dominant-negative constructs coupled to the regulatable promoters into the four test cell lines. Subsequently, the activator is introduced and the repressor, such as rtTA and tTS (see above) via infection using recombinant retroviruses with different selectable markers

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E. Sequences

1. SEQ ID NO:1 Human alpha6 integrin cds (acc# 4557674)

50 atggccgcccgcgggcagctgtgcttgcctctacctgtcggcggggctcctgtcccggtcggcgcagccttcaacttggac
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35 **2. SEQ ID NO:2 Human integrin alpha6 protein sequence (acc# NP_000201):**

1 maaaggqlcll ylsagllsrl gaafnldtre dnvirkygdp gslfgfslam hwqlqpedkr
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 40 181 ghekfsgcqq gvaatftkdf hyivfgapgt ynwkgivrve qknttffdmn ifedgpyevg
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 421 ptqvlkgisp yfgysiagnm dldrnsypdv avgsldsdt ifrsrpvini qktitvtpr
 45 481 idlrqktacg apsgicqlvk scfeytanpa gynpsisivg tleakerrk sglssrvqfr
 541 nqgsepkytq eltlkrqkqk vmeellwlq dnirdklrpi pitasveiqe pssrrrvnsl
 601 pevlpilnsd epktahidvh flkegcgddn vcnslkley kfctregnd kfslpliqkg
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 961 fleysklly ldilmrafid vtaaaenirl pnagtqvrvt vfpsktvaqy sgvpwwiilv
 1021 ailagilmia llvfilwkcg ffrknkdh dyatyhkaeh aqpsdkerlt sda

55 **3. SEQ ID NO:3 Mus musculus alpha6 integrin cds (acc# 7110658):**

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**4. SEQ ID NO:4 Mus musculus alpha6 integrin protein sequence (acc#
 NP_032423)**

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 121 sqgpggkvvt cahryekrqh vntkgesrdi fgrcyvlsgn lrieddmdgg dwsfcdgrlr
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 241 getdhdeslv pvpansylgf sldsgkgivs kdditfvsga pranhsgavv llkrdmksah
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 361 wsnvkpirln gtdksmfgis vknigdingd gypdiavgap yddlqkvfly hgsptgiitk
 421 ptqvlegts p yfgysiagnm dldrnsypdl avgsldsdt ifrsrpvini lktitvtpr
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 961 fleysklly ldilvrasid vtaaaqnikl phagtqrvvt vfpsktvaqy sgvawwiill
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5. SEQ ID NO:5 Human integrin beta4 subunit cds (acc# 6453379):

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6. SEQ ID NO:6 Human beta4 integrin protein sequence (acc# CAB61345):

1 magprpspwa rlllaalisv slsgtlrck kapvksctec vrvdkdcayc tdemfrdrcc
 61 ntqaellaag cqresivvme ssfqiteetq idttlrrsqm spgglrvrlr pgeerhfele
 121 vfeplespvd lyilmdfsns msddldnlkk mgnlarvls qltsdytigf gkfvdkvsvp
 181 qtdmrpeklk epwpnsdppf sfknvislte dvcefrnkliq gerisgnlda peggfdailq
 241 tavctrdigw rpdsthllvf stesafhyea dganvlagim srnderchld ttgtytgyrt
 301 qdypsvptlv rllakhniip ifavtnysys yyeklhityfp vsslgvlqed ssnivellee
 361 afnrirsnlid iraldsprgl rtevtskmfq ktrtgsfhir rgevgiyqvq lralehvdgt
 421 hvcqlpedqk gnihlkpsfs dglkmdagii cdvctcelqk evrsarcsfn gdfvcggcvc
 481 segwsgqtcn cstgslsdiq pclregedkp csgrgecqcg hcvcygegry egqfceydnf

541 qcprtsqfllc ndrgrcsmgq cvcepgwtgp scdcpisnat cidsnggicn grghcecgro
 601 hchqqslytd ticeinysas trasartyap acsarrgapa rrrgarvrna tsrsrwtsl
 661 rearrrwcaa psgtrmttap tatpwkvtap lgptalswct rrrdcpqgsf wwliplllll
 721 lp11a11111 cwkycaccka clallpccnr ghmvqfkdh ymlrenlmas dhldtqmlrs
 5 781 gnlkgrdvvr wkvtnnmqrp gfathaasin ptelvpqgl lrlarlcten llkpdtreca
 841 qlrqveeenl nevyrqisgv hklqqtqkfrq qpnaqkkqdh tivdtvlmap rsakpallkl
 901 tekqvegraf hdlkvapgyy tltadqdarg mvefgegvel vdvrplfir pedddekql
 961 veaidvpagt atlgrrlvni tiikegardv vsfeqpefsv srgdqvarip virrvldcgk
 1021 sqvsyrtqdg taqgnrdyip vegellfqpg eawkelqvl lelqevdsl1 rgrqvrfrhv
 10 1081 qlsnpkfgah lgqphsttii irdpdeldrs ftsqmlssqp pphgd1gapq npnakaagsr
 1141 kihfnwlpss gkpmgyrvky wiqgdsesea hlldskvpsv eltnlypycd yemkvayga
 1201 qgegpysslv scrtthqevps epgrlafnv sstvtqlswa epaetngeit ayevcyglvn
 1261 ddnrpigmk kvldvnpknr mllienlres qpyrytvkar ngagwgpere a1inlatqpk
 1321 rpmsipilpd ipivdaqsge dydsflmysd dvlrspgsq rpsvsddtgc gwkfep1lge
 15 1381 eldlrrvtwr lppeliprls assgrssdae aphgppddgg agkggslpr satpgppgeh
 1441 lvngrmdfaf pgstnslhrm ttttaaaygt hlsphvphrv ltsstl1trd yns1trsehs
 1501 hsttlprdis tltsvsshgl ppiwehgrsr lplswalgsr sraqmkgfpp srgprdsi1l
 1561 agrpaapswg pdsrltagvp dtptrlvfa lgptslrvsw qeprcerplq gysveyqln
 1621 ggellhrlnip npaqtstvve dllpnhsvf rvraqsgew greregvti esqvhpqsp1
 20 1681 cplpgsaftl stpsagplv ftalspds1q lswprrrpn gdivgylvtc emaggggpat
 1741 afrvdgdspe srltvpqlse nvpykfkvqa rttegfger egiitiesqd ggpfpqlgsr
 1801 aglfqhlqs eyssistttht satepflvqp tlgaqhleag gsltrhvtqe fvsrtlttsq
 1861 t1sthmdqf fqt

7. SEQ ID NO:7 Mouse integrin beta4 subunit mRNA (not only cds) (acc# L04678):

25 ggaccgtcgaggcagcgggactgaccagctgggctcactgtattaaagaagcggaccgcgacccggagcggccggggacc
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 caacgcagcccaagtcagagtagtctcactaaggaggaggaggatggcagggccctgttgacagcccatgggtgaagctgc
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8. SEQ ID NO:8 Murine integrin beta4 protein sequence:

MAGPCCSPWV KLLLLARMLS ASLPGDLANR CKKAQVKSCT ECIRVDKSCA YCTDELFKERRCNTQADVLA
 35 AGCRGESILV MESSLEITEN TQIVTSLHRS QVSPQGLQVR LRRGEERTFVFQVFEPLSP VDLIYILMDFS
 NSMSDDLNL KQMGQNLAKI LRQLTSDYTI GFGKFVDKVSVPQTDMRPEK LKEPWPNSDP FFSFKNVISL
 TENVEEFWNK IQGERISGNL DAPEGGFDAIQLTAVCTRI GWRADSTHLL VFSTESAFHY EADGANVLG
 IMNRNDEKCH LDASGAYTQY
 40 KTQDYPVPT LVRLLAHNI IPIFAVTNYS YSYEKLHKY FVSSSLGVLQ EDSSNIVELLEAFYRIRSN
 LDIRALDSPR GLRTEVTSMT LQKTETGSFH IKRGEVGTYN VHLRAVEDIDGTHVCQLAKE DQGGNIHLKP
 SFSDGLRMDA SVICDVCPCE LQKEVRSARC HFRGDFMCGHVCNEGWSGK TCNCSTGSLD DTQPCDLREGE
 DKPCSGHGEC QCGRCVCYGE GRYEGHFCEYDNFQCPRTSG FLCNDRGRCS MGECVCEPGW TCRSCDCPLS
 NATCIDSNGG ICNNGRYCEC
 45 GRCHCNQSL YTDTTCEINY SAILGLCEDL RSCVQCQAWG TGEKKGRACD DCPFKVKMVDLKKKEEVEY
 CSFRDEDDDC TYSYNVEGDG SPGPNSTVLV HKKDCPLAP SWWLIPLLIPLLLLLALLLL LCWKYCACK
 ACLGLLPCCN RGHMVGFKED HYMLRENLMA SDHLDTPMLRSGNLKGRDTV RWKITNNVQR PGFATHAAS
 SPTELVPYGL SLRLGRLCTE NLMKPGTRECDQLRQEVEEN LNEVYRQVSG AHKLQQTFR QCPNAGKKQD
 HTIVDFVLLA PRSAQMLLK
 50 LTEKQVEQGS FHELVKAPGY YTVTAEQDAR GMVEFQEGVE LVDVRVPLFI RPEDDDEKQLLVEAIDVPVS
 TATLGRRLVN ITIIEQASG VVSFEQPEYS VSRGDQVARI PVIRHILDNGKSQVSYSTQD NTAHGHRYV
 PVEGELLFHP GETWKELQVK LLELQEVDSL LRGRQVRRFQVQLSNPKFGA RLQGPSTTTV ILDETDRSLI
 NQTLSSPPPP HGDLGAPQNP NAKAAGSRKIHFNLPPPGK PMGYRVKYWI QGDSESEAH LSKVPSVEL
 TNLYPYCDYE MKVCAYGAKG
 55 EGPYSSLVSC RTHQVPSEP GRLAFNVVSS TVTQLSWAEP AETNGETTAY EVCYGLVNEENRPIGPMKKV
 LVDNPKNRML LIENLRDSQP YRYTVKARNG AGWGPEREAI INLATQPKRPMSPPIIPDIP IVDAQGGEDY
 ENFLMYSDDV LRSPASSQRP SVSDDTEHLV NGRMDFAYPGSANSLHRMTA ANVAYGTHLS PHLSHRVLST
 SSTITRDYHS LTRTEHSHSG TLPRDYSTLTLSSQASLLS GKMGGAGFRC PGLLGP

9. SEQ ID NO:9 Mouse beta4 dominant negative cds:

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 60 gccaacgctgcaagaaggctcaggtgaagagctgtaccgagtgcatccgggtggacaagagctgtgcctactgcacagac
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10. SEQ ID NO:10 Mouse beta4 dominant negative protein sequence:

MAGPCCSPWV KLLLLARMLS ASLPGDLANR CKKAQVKSCT ECIRVDKSCA YCTDELFKERRCNTQADVLA
 AGCRGESILV MESSLEITEN TQIVTSLHRS QVSPQGLQVR LRRGEERTFVFQVFEPLSP VDLIILMDFS
 25 NSMSDDLNL KMQGNLAKI LRQLTSDYTI GFGKFVDKVSVPQTDMRPEK LKEPWPNSDP PFSFKNVISL
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 IMNRNDEKCH LDASGAYTQY
 KTQDYPSPVT LVRLLAKHNI IPIFAVTNYS YSYYEKLHKY FPFVSSLGLVQ EDSSNIVELLEAFYRIRSN
 LDIRALDSPR GLRTEVTSNT LQKTETGSFH IKRGEVGTYN VHLRAVEDIDGTHVCQLAKE DQGGNIHLKP
 30 SFSDGLRMDA SVICDVCPCE LQKEVRSARC HFRGDFMCGHCVCNNEGWSGK TCNCSTGSLD DTQPCLEGE
 DKPCSGHGEC QCGRCVCYGE GRYEGHFCEYDNFQCPRTSG FLCNDRGRCS MGECVCEPGW TGRSCDCPLS
 NATCIDSNGG ICNNGRGYCEC
 GRCHCNQQLS YDITTCBINY SAILGLCEDL RSCVQCQAWG TGEKKGRACD DCPFKVKMVDELKKEEVVEY
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11. SEQ ID NO:11 Murine laminin gamma2 chain complete cds (acc# U43327):

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 caggctgagcaggcccttcaggacattctgggagaagctcagatttcagaaggggcaatgagagccgttgctg
 tccggctggccaaggcaaggagccaagagaacgactacaagaccccgctggatgacctcaagatgactgcaga
 aaggatccgggcccctgggagtcagcatcagaacagagttcaggatacgagcagactcatctctcagatgcgc
 5 ctgagtctggcaggaagcgaagctctcttggaaaacactaatatccattctcttgagcactacgtggggccga
 atgattttaaaagtctggctcaggaggctacaagaaaggcagacagccacgctgagtcagctaacgcaatgaa
 gcaactagcaagggaactgaggactactccaaacaagcacttttcattggcccgcaagctcttgagtggagga
 ggcggaagtggctcttgggacagctccgtgggtacaaggctcttatgggaaaattagagaaaaccaagtccctga
 gccagcagctgtcattggaggggcacccaagccgacabtgaaagctgataggctcgatcagcacagctctccgcct
 10 cctggattctgctctcagcttcagggagtcagtgatctgtcctttcaggtggaagcaaagaggatcagacaa
 aaggctgattctctctcaaacctgggtgaccagacaaacggatgcattcacgcgtgtgcgaaacaatctgggga
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 aacatcctgaagaacctccgagagtttgatctgcagggtgaagacagaaaagcagaggctgaagaggccatga
 15 agagactctcctctattagccagaagggttgaggatgccagtgacaagacccagcaagcagaaaacggccctggg
 gagcgccactgcccagacaccaacgggcaaagaacgcagctagggaggccctggagatcagcagcgagatagag
 ctggagatagggagctctgaacttggaagctaattgtgacagcagatggggcccttgggcatggagaaagggactg
 ccactctgaagagcgagatgagagagatgattgagctggccagaaaggagctggagtttgacacggataagga
 cacgggtgcagctgggtgattactgaagcccagcaagctgatgccagagccacagagtgccggagttaccatccaa
 20 gacacrcctcaacacattgggacggcatcctacacctcatagaccagcctggcagtggtgatgaagaagggatga
 tgctattagaacaagggtttttccaagccaagaaccagatcaacagtcgacttcggcccttgatgtctgacct
 ggaggagagggtgcgtcggcagaggaaccacctccatctgctggagactagcatagatggaattcttgctgat
 gtgaagaacctggagaacattcgagacaacctgccccaggctgctacaataaccaagctcttgagcaacagt
 ga

25 **12. SEQ ID NO:12 Murine laminin gamma2 chain protein sequence (acc#
 AAA85256)**

1 mpalwlsccl gvallllpasq atsrrevcdc ngksrqcvfd qelhrqagsg frclncndnt
 61 agvhcersre gfyqhqsksr clpncnchskg slsagcdnsg qcrckpgvtg qrcdqccpgf
 121 hmltdagctr dqgqldskcd cdpagisgpc dsgrcvckpa vtgercdrcr prdyhldran
 30 181 pegctqcfcy ghsaschasa dfsvhkitst fsqdvdgwka vqrngapakl hwsqrhrdvf
 241 ssarrsdpyv fvapakflgn qqvsyqgsls fdyrvdrggr qpsaydvile gaglqirapl
 301 mapgkltlpcg itktytfrln ehpsshwspq lsyfeyrrll rnltallmir atygeystgy
 361 idnvtlvsar pvlgapapwv ercvcllgyk gqfcqecag ykrdsarlga fgacvpcncq
 421 gegacdptg dcysgdenpd iecadcpigf yndphdprsc kpcpchnqfs csvmpeteev
 35 481 vcnnpppgvt garcelcadg ffgdppfgeh pvrpcqrcq nnnvdpnasg ncdqltgrcl
 541 kciyntagvy cdqckagyfy dplapnpadk crancspmg aepgecrdg scvckpgfga
 601 fncdhaalts cpacynqvki qmdqftqqlq slealvskaq ggggggtvpv qlegrieqae
 661 qalqdlgea qisegamrav avrlakarsq endyktrldd lkmtaerira lgsqhqnrvq
 721 dtsrlisqmr lslagseall entnihsseh yvgpndfksl aqeatrkads haesanamkq
 40 781 laretedysk qalslarkll sggsgsgswd ssvvqglmkg lektkslsqg lslegtqadi
 841 eadrsyqhs1 rlldsasqlg gvsdlsfqve akrirqkads lsnlvtrqtd aftrvrrnlq
 901 nweketrqll qtgkdrqrts dqlsranla knragealsm gnatfyeven ilknlnrefdl
 961 qvedrkaeae eamkrllsis qkvadasdkt qqaetalgsa tadtqrakna arealeisse
 1021 ieleigslnl earvntadgal amekgtatl semremiela rkelefddtdk dtvqlvitea
 45 1081 qqadaratsa gvtiqdtlnt ldgilhlidq pgsvdeegmm lleqglfqak tqinsrlrpl
 1141 msdleervrr qrnhlhllet sidgiladvk nlenirdnlp pgcyntqale qq

**13. SEQ ID NO:13 Human laminin gamma2 chain complete cds (acc#
 AH006634)**

atgcctgcgctctggctgggctgctgcctctgcttctcgctcctcctgcccgcagcccgggccacctccaggaggggaagtc
 50 tgtgattgcaatgggaagtccaggcagtgatctttgatcggggaacttcacagacaaactggtaatggattccgctgcctc
 aactgcaatgacaacactgatggcattcactgcgagaagtgcagaatggcttttaccggcacagagaaagggaccgctgt
 ttgccctgcaattgtaactccaaagggttctcttagtgctcgatgtgacaactctggacgggtgcagctgtaaaccagggtgtg
 acaggagccagatgcgaccgatgtctgccaggcttccacatgctcacggatgcgggggtgcacccaagaccagagactgcta
 gactccaagtgtgactgtgacccagctggcatcgcaggggccctgtgacgcgggcccgtgtgtctgcaagccagctgttact
 55 ggagaacgctgtgatagggtgtcgatcaggttactataacttggtatggggggaaacctgagggtgttaccagtgcttctgc
 tatgggcatccagccagctgcgcagctctgcagaatacagtgctccataagatcacctctacctttcatcaagatgttgat
 ggctggaaggctgtccaacgaaatgggtctcctgcaaagctccaatgggtcacagcgcctatcaagatgtgttttagctcagcc
 caacgactagacctgtctatcttgggtcctgcccatttcttgggaatcaacagggtgagctatgggcaaagcctgtcc
 tttgactaccgtgtggacagaggaggcagacacccatctgcccattgatgtgattctggaagggtgctgggtctacggatcaca
 60 gctcccttgatgccacttggaagacactgccttggtgggtcaccaagacttacacattcagggttaaatgagcatccaagc
 aataattggagccccagctgagttactttgagtatcgaagggttactgcgggaatctcacagccctccgcatccgagctaca
 tatggagaatacagtaactgggtacattgacaatgtgacctgatttcagccccctgtctctggagccccagcaccctgg

gttgaacagtgatatgtcctgttgggtacaaggggcaattctgccaggattctgtcttctgggtacaagagagattcagcg
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 tcaggggatgagaatcctgacattgagtgctgactgcccaattgggttctacaacgatccgcacgacccccgcagctgc
 aagccatgtccctgtcataacgggttcagctgctcagtgatgccggagacggaggaggtggtgtgcaataactgccctccc
 5 ggggtcaccggtgcccgtgtgagctctgtgctgatggctactttggggaccctttgggtgaacatggcccagtgaggcct
 tgtcagccctgtcaatgcaacaacaatgtggaccocagtgccctctgggaattgtgaccggctgacaggcaggtgtttgaag
 tgtatccacaacacagccggcatctactgcgaccagtgcaagcaggctacttcggggaccattgggtcccaaccagca
 gacaagtgtcgagcttgcaactgtaaccccatgggctcagagcctgtaggatctcgaagtgtggcacctgtgtttgcaag
 ccaggattttgggtggccccaaactgtgagcatggagcattcagctgtccagcttctataatcaagtgaagattcagatggat
 10 cagtttatgcagcagcttcagagaatggaggccctgatttcaaaggctcaggggtggtgatggagtagtacctgatacagag
 ctggaaggcagatgcagcaggctgagcaggcccttcaggacattctgagagatgccagatttcagaagggtgctagcaga
 tcccttgggtctccagttggccaagggtgaggagccaagagaacagctaccagagccgctggatgacctcaagatgactgtg
 gaaagagttcgggctctgggaagttagtaccagaaccgagttcgggatactcacaggctcatcactcagatgcagctgagc
 ctggcagaaaagtgaagcttcccttgggaaacactaacaattcctgcctcagaccactacgtggggccaaatggctttaaaagt
 15 ctggctcaggaggccacaagatttagcagaaagccacgttgagttagccagtaacatggagcaactgacaagggaaactgag
 gactattccaaacaagccctctcactgggtgcgcaaggccctgcagtaaggagtcggaagcggagcggtagcccgagcgg
 gctgtggtgcaagggcttgtggaaaaattggagaaaaccaagtccttggccagcagttgacaagggaggccactcaagcg
 gaaattgaagcagatagggtcttatcagcacagctctccgctcctggattcagtgctctcggcttcagggagtcagtgatcag
 tcccttccaggtggaagaagcaaaaggatcaaaacaaaagcggattcactctcaacgctggtaaccaggcatatggatgag
 20 ttcaagcgtacacaaaagaatctgggaaactggaaagaagaagcacagcagctcttacagaatggaaaaagtgggagagag
 aaatcagatcagctgctttcccgctgccaatcttgctaaaagcagagcacagaagcactgagtatgggcaatgccactttt
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 atgaagagactctcctacatcagccagaagggtttcagatgccagtgacaagaccagcaagcagaaagagccctggggagc
 gctgctgctgatgcacagagggcaaagaatggggccggggaggccctggaaatctccagtgagattgaacaggagattggg
 25 agtctgaacttggaaagccaatgtgacagcagatggagccttggccatggaaaagggactggcctctctgaagagtgaagt
 agggaaagtggaaaggagagctggaaaggaaggagctggagtttgacacgaatatggatgcagtacagatggtgattacagaa
 gccagaagggttgataccagagccaagaacgctgggggtacaatccaagacacactcaacacattagacggcctcctgcac
 ctgatggaccagcctctcagtgtagatgaagaggggctggtcttactggagcagaagctttcccgagccaagaccagatc
 aacagccaactcgggcccattgatgtcagagctggaagagagggcacgtcagcagagggggccacctccatttgctggagaca
 30 agcatagatgggattctggctgatgtgaagaacttggagaacattagggacaacctgccccaggctgctacaataccag
 gctcttgagcaacagtga

14. SEQ ID NO:14 Human laminin gamma2 chain protein sequence (acc# AAC50457) Alternative splice form

1 mpalwlgccl cfslllpaar atsrrevcdc ngksrqcifd relhrqtgng frclncndnt
 35 61 dgihecekcn gfyrrhrerdr clpcncnsgk slsarcnsg rscckpgvtg arcdrclpgf
 121 hmltdagctq dqrllldskcd cdpagiagpc dagrcvckpa vtgercdrcr sgyynldggn
 181 pegctqcfcy ghsascrssa eysvokitst fhqdvdgwka vqrngspakl qwsqrhgdvf
 241 ssagrldpvy fvapakflgn qqvsyqsls fdyrvdrgr hpsahdvile gaglritapl
 301 mplgkltlpcg ltktytfrln ehpsnnwspq lsyfeyrll rnlalrira tygeystgyi
 40 361 dnvltlisarp vsgapapwve qcicpvgykg qfcqdcasgy krdsarlpgf gtcipncqg
 421 ggacdptgd cysgdenpdi ecadcpigfy ndphdprsc pcpchngfsc svmpeteevv
 481 cncpppgvtg arcelcadgy fgdpfgehgp vrpcqpcqcn nnvdpsasgn cdrltgrclk
 541 cihntagiyc dqckagyfgd plapnpadkc racncnmpgs epvgcrsdgt cvckpgfggp
 601 ncehgafscp acynqvkiqm dqfmqqlqrm ealiskaagg dgvpdtele grmqqaegal
 45 661 qdlrdaqis egasrslglq lakvrsqens yqsrlddlkm tvervralgs qyqnrvrth
 721 rlitmqqlsl aeseaslgnt nipasdhvyg pngfkslage atrlaeshve sasnmeqltr
 781 etedyskqal slvrkalheg vsgsgspdg avvgglvekl ektkslaqql treatgaeie
 841 adrsyqhsrlr lldsvsrlqg vsdqsfqvee akrikqkads lstlvtrhmd efkrtqknlg
 901 nwkeeaqql qngksgreks dqllsranla ksraqealsm gnatfyeves ilknlrefdl
 50 961 qvdnrkaeae eamkrlsyis qkvsdasdkt qqaeralgsa aadaqrakng agealeisse
 1021 ieqeigslnl eanvtadgal amekglaslk semrevegele erkelefdtn mdavqmvite
 1081 agkvdtrakn agvtiqdtln tldgllhlmd qplsvdeegl vlleqklsla ktqinsqlrp
 1141 mmseleerar qqrghlhlle tsidgiladv knlenirdnl ppgcyntgal eqq

15. SEQ ID NO:15 Human laminin gamma2 chain protein sequence (acc# AAC50456) Alternative splice form

1 mpalwlgccl cfslllpaar atsrrevcdc ngksrqcifd relhrqtgng frclncndnt
 61 dgihecekcn gfyrrhrerdr clpcncnsgk slsarcnsg rscckpgvtg arcdrclpgf
 121 hmltdagctq dqrllldskcd cdpagiagpc dagrcvckpa vtgercdrcr sgyynldggn
 181 pegctqcfcy ghsascrssa eysvokitst fhqdvdgwka vqrngspakl qwsqrhgdvf
 60 241 ssagrldpvy fvapakflgn qqvsyqsls fdyrvdrgr hpsahdvile gaglritapl
 301 mplgkltlpcg ltktytfrln ehpsnnwspq lsyfeyrll rnlalrira tygeystgyi
 361 dnvltlisarp vsgapapwve qcicpvgykg qfcqdcasgy krdsarlpgf gtcipncqg
 421 ggacdptgd cysgdenpdi ecadcpigfy ndphdprsc pcpchngfsc svmpeteevv
 481 cncpppgvtg arcelcadgy fgdpfgehgp vrpcqpcqcn nnvdpsasgn cdrltgrclk
 65 541 cihntagiyc dqckagyfgd plapnpadkc racncnmpgs epvgcrsdgt cvckpgfggp

10

10

10

acacgggactacaactcactgacccgctcagaacactcacactcgaccacactgcccagggaactactccaccctcacctcc
 gtctcctcccacggcctccctcccatcttggaacacgggaggagcaggcttccgctgtcctgggncctggggtcccggagt
 cgggctcagatgaaagggttcccccttccaggggcccacgagactctataatcctggctgggaggccagcagcgcctcc
 tggggcccagactctcgctgactgctggtgtgcccgaacgcccacccgctggtgttctctgcccctggggcccacatct
 5 ctcagagtgaagctggcaggagccgcggtgagcgaggccgctgagggctacagtgtgagtaaccagctgctgaacggcggg
 gagctgcatcggtcaacatcccccaaccctgcccagacctcggtggtggtggaagacctcctgcccaccactcctacgtg
 ttccgctgcccgggcccagagccaggaaggctggggcccagagcgtgaggggtgtcatcaccattgaatcccagggtgcacccg
 cagagcccactgtgtccccctgcccaggctccgcttctactttgagcactcccagtgccccaggcccgctggtgttactgccc
 ctgagcccagactcgctgagctgagctgggagcggccacggaggcccaatggggatctcgtcgggtacctggtgacctgt
 10 gagatggcccaggaggaggtccagccaccgcatctccgggtggtgagagacagcccagagagccgggtgacctgcccgggc
 ctcagcgagaacgtgcccctacaagttcaagggtgagggccaggaccactgagggcttccgggcccagagcgcgaggggcatc
 accatagagtcccaggatggaggtcccttcccgacgtgggcagccgtgcccgggtctctccagcaccgctgcaaagcgag
 tacagcagcatctccaccaccacaccagcggccaccgaccccttctagtgggtccgaccctggggggcccagcactggag
 gcaggcggtccctcaccggcatgtgaccaggagtttgtgagccggacactgaccaccagcgggaacccttagcaccac
 15 atggaccaacagttcttccaaacttga

17. SEQ ID NO:17 Human beta4 integrin protein sequence Variant V at 34 to I:

1 magprpwpwa rlllaalisv slsgtlnrck kaplksctec vrvdkdcayc tдемfrdrcc
 61 ntqaellaag cqresivvme ssfqiteetq idttlrrsqm spqglrvrlr pgeerhfele
 20 121 vfeplespvd lyilmdfsns msddldnlkk mggnlarvls qltsdytigf gkfvdksvsv
 181 qtdmrpeklk epwpnsdppf sfknvislte dvdefrnklq gerisgnlda peggfdailq
 241 tavctrdigw rpdsthllvf stesafhyea dganvlagim srnderchld ttgtytqyrt
 301 qdypsvptlv rllakhniip ifavtnsys yyeklhtyfp vsslgvlqed ssnivellee
 361 afnrirsnd iraldsprgl rtevtskmfq ktrtgsfhir rgevgyqvq lralehvdgt
 25 421 hvcqlpedqk gnihlkpsfs dglkmdagii cdvctcelqk evrsarcsfn gdfvcggcvc
 481 segwsgqtcn cstgslsdiq pclregedkp csgrgecqcq hcvcygegry eggficeydnf
 541 qcprtsqflc ndrgrcsmgq cvcepwtgtp scdcpnsnat cidsgngicn grghcecgrc
 601 hchqqslytd ticeinysas trasartyap acsarrgapa rrrgarvrna tsrsrwwtsl
 661 rearrrwcaa psgtrmttap tatpwkvtp lgptalswct rrrdcpqgsf wwliplllll
 30 721 lpllllllll cwkycaccka clallpcnr ghmvfgkedh ymlrenlmas dhldtpmlrs
 781 gnlkgrdvvr wkvtnmqrp gfathaasin ptelvpysls lrlarlcten likpdtreca
 841 qlrqveenl nevyrgisgv hklqgtkfrq qpnaqkkqdh tivdtvlmap rsakpallkl
 901 tekqvegraf hdlkvapgyy tltadqdarg mvefgegvel vdvrvplfir pedddekqll
 961 veaidvpagt atlgrrlvni tiikeqardv vsfeqpefsv srgdqvarip virrvldggk
 35 1021 sqvsyrtqdg taqgnrdyip vegellfqpg eawkelqvkl lelgevdsll rgrqvrfrhv
 1081 qlsnpkfgah lgqphstti iirdpeldrs ftsqmlssqp pphgdilgapq npnakaagsr
 1141 kihfnwlpss gkpmgyrvky wiqgdsesea hllldskvpsv eltnlypycd yemkvcaiga
 1201 qgegpysslv scrthqevps epgrlafnv sstvtqlswa epaetngeit ayevcyglvn
 1261 ddnrpigmk kvlvdpknr mllienlres qpyrytvkar ngagwgpere aiinlatqpk
 40 1321 rpmsipiipd ipivdaqsge dydsflmysd dvlrspsgsq rpsvsddtgc gwkfepllge
 1381 eldlrrvtwr lppeliprls assgrssdae aphgppddgg aggkgsllpr satpgppgeh
 1441 lvngrmdfaf pgstnslhrm tttssaaaygt hlsphvphrv ltsstlttrd ynsltrsehs
 1501 hsttlprdis tltsvsshgl ppiwehgrsr lplswalgsr sraqmkgfpp srgprdsiil
 1561 agrpaapswg pdsrltagvp dtptrlvfa lgptslrvsw qeprcerplq gysveyqln
 45 1621 ggelhrlnip npqtsvvve dlipnhsyvf rvraqsqegw greregviti esqvhpspl
 1681 cplpgsaftl stpsagplv ftalspdsq lswerprprn gdivgylvtc emaggggpat
 1741 afrvdgspe srltvpglse nvpykfkva rtteggfper egiitiesqd ggpfpqlgsr
 1801 aglfqkplqs eyssistttht satepflvvp tlgaghleag gsltrhvtqe fvsrtlttsq
 1861 tlsthmdqqf fqt

18. SEQ ID NO:18 Human integrin beta4 subunit cds Variant at 34 V to I:

atg gca ggg cca cgc ccc agc cca tgg gcc agg ctg ctc ctg gca gcc ttg atc agc gtc
 agc ctc tct ggg acc ttg aac cgc tgc aag aag gcc cca AtA aag agc tgc acg gag tgt
 gtc cgt gtg gat aag gac tgc gcc tac tgc aca gac gag atg ttc agg
 gaccggcgctgcaacacccaggcggagctgctggccgcgggctgcccagcgggagagcatcgtggtcatggagagcagcttc
 55 caaatcacagaggagacccagattgacaccaccctgcggcgagccagatgtcccccaaggcctgcgggtccgtctgcgg
 cccggtgaggagcggcattttgagctggaggtgtttgagccactggagagccccgtggacctgtacatcctcatggacttc
 tccaactccatgtccgatgatctggacaacctcaagaagatggggcagaacctggctcgggtcctgagccagctcaccagc
 gactacactattggatttggaagtttgggacaaagtcagcgtcccgcagacggacatgaggcctcagaagctgaaggag
 ccttggcccacagtgaccccccttctccttcaagaacgtcatcagcctgacagaagatgtggatcagttccggaataaa
 60 ctgcaggagagcggatctcaggcaacctggatgctcctgaggcggtctcgatgccatcctgcagacagctgtgtgcacg
 agggacattggctggcgcccgagcagcaccacctgctggtcttctccaccagctcagccttccactatgaggctgatggc
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 tactcctatagctactacgagaagcttcacacctatttccctgtctcctcactgggggtgctgcaggaggactcgtccaac
 65 atcgtggagctgctggaggaggccttcaatcggatccgctccaacctggacatccgggcccctagacagccccgaggcctt

cggacagaggtcacctccaagatgttccagaagacgaggactgggtcctttcacatccggcggggggaagtgggtatatac
 caggtgcagctgcgggccccttgagcacgtggatgggacgcacgtgtgccagctgccggaggaccagaagggcaacatccat
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 5 tgcaactgctccaccggctctctgagtgacattcagccctgcctgcgggagggcgaggacaagccgtgctccggcgggtggg
 gagtgccagtgcgggcactgtgtgtgtgtacggcgaaggccgtacgagggtcagtctctgcgagtatgacaacttccagtgt
 ccccgcaacttccgggttccctctgcaatgaccgaggacgtgctccatggggccagtgtgtgtgtgtgagcctgggtggacaggc
 ccaagctgtgactgtccctcagcaatgccacctgcctgcagcaatgggggcatctgtaatggacgtggccactgtgag
 tgtggcgctgacctgccaccagcagtcgctctacacggacaccatctgcgagatcaactactcggcgctccaccgggccc
 10 tctgcgaggacctacgctcctgcgtgcagtgcaggcggtggggcaccggcgagaagaagggcgccagctgtgaggaatgca
 acttcaaggtcaagatgggtggacgagcttaagagaggcgaggaggtgggtgggtgcgctgctccttccgggacgaggatgacg
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 15 gaccactacatgctgcgggagaacctgatggcctctgaccaacttggaacgcgccatgctgcgcagcgaggaaacctcaagggc
 cgtgacgtgggtccgctggaaggtcaccaacaacatgcagcgccctggcctttgccactcatgcgcagcatcaaccccaca
 gagctgggtgccctacgggctgtccttgccgctggcccgccctttgcacgcgagaacctgctgaagcctgacactcgggagtgc
 gccagctgcgcagaggaggtggaggagaacctgaacgaggtctacaggcagatctccgggtgtacacaagctccagcagacc
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19. SEQ ID NO:19 Degenerate Human integrin beta4 subunit cds variant at 34 Vto 1:

atg gca ggg cca cgG ccc agc cca tgg gcc agg ctg etc ctg gca gcc ttg atc agc gtc
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What is claimed is:

1. A method of reducing the proliferation of a cancer cell which comprises inhibiting ligand binding to an integrin receptor on the cancer cell, wherein the integrin receptor comprises an integrin.
2. The method of claim 1, wherein the integrin receptor comprises integrin B4.
3. The method of claim 1, wherein the integrin receptor comprises integrin A6.
4. The method of claim 1, wherein the ligand that binds to the integrin receptor is laminin5.
5. The method of claim 1, wherein the integrin receptor is A6B4.
6. The method of claim 1, wherein the ligand comprises laminin5.
7. The method of claim 1, wherein the ligand comprises the gamma-2 subunit of laminin5.
8. The method of claim 1, wherein inhibiting ligand binding to an integrin receptor comprises contacting a A6 integrin with a composition that inhibits ligand binding.
9. The method of claim 1, wherein inhibiting ligand binding to an integrin receptor comprises contacting a B4 integrin with a composition that inhibits ligand binding.
10. The method of claim 1, wherein inhibiting ligand binding to an integrin receptor comprises contacting a laminin5 with a composition that inhibits ligand binding.
11. The method of claim 1, wherein inhibiting ligand binding to an integrin receptor comprises contacting a gamma-2 subunit with a composition that inhibits ligand binding.
12. A method of reducing the proliferation of a cancer cell which comprises reducing integrin-integrin interaction, integrin receptor clustering interaction, or integrin-non-integrin protein interaction.
13. The method of claim 12, wherein reducing integrin-integrin interaction, integrin receptor clustering interaction, or integrin-non-integrin protein interaction comprises contacting a A6 integrin with a composition that inhibits an interaction between the B4 integrin and another integrin or protein molecule.
14. The method of claim 12, wherein reducing integrin-integrin interaction, integrin receptor clustering interaction, or integrin-non-integrin protein interaction comprises contacting a B4 integrin with a composition that inhibits the interaction between the B4 integrin and another integrin or protein molecule..
15. The method of claim 12, wherein reducing integrin-integrin interaction, integrin receptor clustering interaction, or integrin-non-integrin protein interaction comprises contacting a

laminin5 with a composition that inhibits ligand binding.

16. The method of claim 12, wherein reducing integrin-integrin interaction, integrin receptor clustering interaction, or integrin-non-integrin protein interaction comprises contacting a gamma2 subunit with a composition that inhibits ligand binding.

17. The method of claim 12, wherein the non-integrin protein comprises a growth factor receptor.

18. The method of claim 12, wherein the non-integrin protein comprises a hemidesmosome junction.

19. The method of claim 12, wherein the non-integrin protein comprises a SH2 domain.

20. The method of claim 19, wherein the non-integrin protein comprises a Shc protein.

21. The method of claim 12, wherein the non-integrin protein comprises a IRS-1 protein.

22. The method of claim 12, wherein the non-integrin protein comprises a IRS-2 protein.

23. A method of reducing the proliferation of a cancer cell which comprises reducing the production of an integrin by the cancer cell.

24. The method of claim 23, wherein the production of an integrin is reduced by inhibiting signaling leading to induction of expression of an integrin.

25. The method of claim 23, wherein reducing the production of an integrin comprises inhibiting alpha6 production.

26. The method of claim 25, wherein inhibiting alpha6 production further comprises using antisense molecules to alpha6 mRNA.

27. The method of claim 23, wherein reducing the production of an integrin comprises inhibiting beta4 production.

28. The method of claim 27, wherein inhibiting beta4 production further comprises using antisense molecules to beta4 mRNA.

29. A method of reducing the proliferation of a cancer cell which comprises reducing the production of an integrin receptor ligand by the cancer cell.

30. The method of claim 29, wherein reducing the production of an integrin receptor ligand comprises inhibiting gamma2 production.

31. The method of claim 29, wherein reducing the production of an integrin receptor ligand comprises inhibiting laminin production.

32. The method of claim 29, wherein reducing the production of an integrin receptor

ligand comprises inhibiting laminin5 production.

33. A method of reducing the proliferation of a cancer cell which comprises interfering with an integrin signaling pathway.

34. The method of claim 33, wherein interfering with an integrin signaling pathway comprises contacting an A6 integrin in the cell with a composition that inhibits ligand binding.

35. The method of claim 33, wherein interfering with an integrin signaling pathway comprises contacting a B4 integrin with a composition that inhibits ligand binding.

36. The method of claim 33, wherein interfering with an integrin signaling pathway comprises contacting a laminin5 with a composition that inhibits ligand binding.

37. The method of claim 33, wherein interfering with an integrin signaling pathway comprises contacting a gamma2 subunit with a composition that inhibits ligand binding.

38. The method of claim 33, wherein interfering with an integrin signaling pathway comprises contacting the cancer cell with a molecule that interferes with at least one of talin, paxillin, vinculin, a CAS family protein, CRX, NCK, FAK, ILK, Src, Fyn, Shc, Grb-2, Guanine nucleotide exchange factors, SOS, DOCK 180, Vav, Syk, P-1-3 kinase, AKT, Bad, Bid, Caspase 9, Cdc42, PAK, Rac, Rho, Rho kinase, Ras, Caveolin, Tetraspan, Receptor-type protein tyrosine phosphatase, SHP-2, Alpha-actinin, Filamin, Cytohesin, Beta3-endonexin, ICAP-1, RACK-1, CIB, actin, receptor tyrosine kinase, IRS-1 or IRS-2.

39. The method of claim 33, wherein interfering with an integrin signaling pathway comprises contacting the cancer cell with an agent that interferes with post-translational modification of integrins.

40. The method of claim 39, wherein the post translational modification is glycosylation or phosphorylation.

41. The method of claims 1, 12, 23, 29, 33, wherein the integrin comprises an A6 integrin.

42. The method of claims 1, 12, 23, 29, 33, wherein the integrin comprises a B4 integrin.

43. The method of claims 1, 12, 23, 29, 33, further comprising reducing a laminin5-integrin interaction.

44. The method of claims 1, 12, 23, 29, 33, further comprising reducing a laminin5 gamma2 integrin interaction.

45. The method of claims 1, 12, 23, 29, 33, wherein reducing the proliferation of the cancer cells is selective.

46. The method of claims 1, 12, 23, 29, 33, wherein the cancer cell is not an MDA-MB-

435 cell.

47. The method of claims 1, 12, 23, 29, 33, wherein the cancer cell is not an HMT-3522 cell.

48. The method of claims 1, 12, 23, 29, 33, wherein the cancer cell is not an RKO colon carcinoma line.

49. The method of claims 1, 12, 23, 29, 33, wherein the cancer cell does not express exogenous B4 integrin.

50. The method of claims 1, 12, 23, 29, 33, further comprising contacting the cancer cells with a small molecule, peptide, peptide mimetic, or oligonucleotide or synthetic analog thereof.

51. The method of claim 50, wherein the cancer cells are contacted with dominant-negative beta 4 integrin.

52. The method of claim 50, wherein the cancer cell is contacted with an antisense molecule.

53. The method of claim 52, wherein the antisense molecule is linked to a leader sequence which enables translocation across a cell membrane.

54. The method of claim 53, wherein the leader sequence binds to a cell surface protein which facilitates internalization.

55. The method of claim 54, wherein the leader sequence is TAT or antennapedia, or fragment thereof.

56. The method of claim 52, wherein the antisense molecule is an alpha6 RNA antisense molecule.

57. The method of claim 47, wherein the small molecule peptide, peptide mimetic, or oligonucleotide or synthetic analog thereof is linked to a carrier.

58. The method of claim 57, wherein the carrier is at least one of a lipidic carrier, charged carrier, retroviral carrier, TAT or fragment thereof, antennapedia or fragment thereof, or polyethylene glycol.

59. The method of any one of claims 1, 12, 23, 29, 33, further comprising contacting the cancer cell with another agent which modulates cell signaling, a chemotherapeutic drug, or treated with radiation or angiogenesis inhibitor.

60. The method of any of claims 1, 12, 23, 29, 33, wherein reducing the proliferation of cancer cell is *in vitro*.

61. The method of any of claims 1, 12, 23, 29, 33, wherein reducing the proliferation

of the cancer cell is *in vivo*.

62. The method of any of claims 1, 12, 23, 29, 33, wherein the cancer cell is selected from the group consisting of melanoma, adenoma, lymphoma, myeloma, carcinoma, plasmocytoma, sarcoma, glioma, thymoma, leukemia, skin cancer, retinal cancer, breast cancer, prostate cancer, colon cancer, esophageal cancer, stomach cancer, pancreas cancer, brain tumors, lung cancer, ovarian cancer, cervical cancer, hepatic cancer, gastrointestinal cancer, and head and neck cancer cells.

63. The method of claims 1, 12, 23, 29, 33, wherein the cancer cell is killed.

64. The method of claims 1, 12, 23, 29, 33, wherein the cancer cell expresses a mutated Ras.

65. The method of claims 1, 12, 23, 29, 33, wherein the cancer cell expresses a mutated Ras and a mutated p53.

66. The method of claims 1, 12, 23, 29, 33, wherein the cancer cell expresses a mutated Ras and activates the AKT/PKB protein.

67. The method of claims 1, 12, 23, 29, 33, wherein the cancer cell expresses a mutated Ras, a mutated p53, and activates the AKT/PKB protein.

68. The method of claims 1, 12, 23, 29, 33, wherein the cancer cell expresses a mutated APC.

69. The method of claims 1, 12, 23, 29, 33, wherein the cancer cell expresses a mutated Ras and mutated APC.

70. A method of isolating molecules that bind with a target molecule selected from the group consisting, B4 integrin, alpha6 integrin, and the gamma2 subunit of laminin5 comprising 1) contacting a library of molecules with the target molecule and 2) collecting molecules that bind the target molecule producing an enriched population of molecules.

71. The method of claim 70, further comprising the step of repeating steps 1 and 2 with the enriched population of molecules.

72. The method of claim 70, wherein the library comprises a small molecule, peptide, peptide mimetic, or oligonucleotide.

73. A method of assessing a subject's risk of developing cancer comprising determining the amount of A6 present in a target cell obtained from the subject, wherein a determination of increased levels of A6 correlates with an increased risk of cancer.

74. The method of claim 73, further comprising comparing the amount A6 present to the

amount in a control cell.

75. The method of claim 73, wherein determining the amount of A6 present in the target cell comprises assaying the amount of A6 mRNA in the cell.

76. The method of claim 75, wherein the assaying the amount of mRNA in the cell comprises hybridizing a A6 probe to a sample of the subject's mRNA.

77. The method claim 75, wherein the assaying the amount of mRNA in the cell comprises hybridizing a A6 primer to a sample of the subject's mRNA.

78. The method claim 77, wherein the assaying the amount of mRNA further comprises performing an nucleic acid amplification reaction involving the primer.

79. The method of claim 78, wherein the nucleic acid amplification reaction comprises reverse transcription, producing a cDNA.

80. The method of claim 79, wherein the nucleic acid amplification reaction further comprises performing a polymerase chain reaction on the cDNA.

81. The method of claim 73, wherein determining the amount of A6 present in the target cell comprises assaying the amount of A6 protein in the cell.

82. A method of assessing a subject's risk of acquiring cancer comprising determining the amount of B4 present in a target cell obtained from the subject, wherein a determination of increased levels of B4 correlates with an increased risk of cancer.

83. The method of claim 82, further comprising comparing the amount B4 present to the amount in a control cell.

84. A method of assessing a subject's risk of acquiring cancer comprising determining the amount of laminin5 present in a target cell obtained from the subject, wherein a determination of increased levels of laminin5 correlates with an increased risk of cancer.

85. The method of claim 84, further comprising comparing the amount laminin5 present to the amount in a control cell.

86. A method of assessing a subject's risk of acquiring cancer comprising determining the amount of gamma2 subunit present in a target cell obtained from the subject, wherein a determination of increased levels of gamma2 subunit correlates with an increased risk of cancer.

87. The method of claim 86, further comprising comparing the amount gamma2 subunit present to the amount in a control cell.

88. A method of reducing the proliferation of a cancer cell in a patient which comprises administering to the patient a composition which inhibits ligand binding to an integrin receptor on

the cancer cell, wherein the integrin receptor comprises an integrin.

89. A method of reducing the proliferation of a cancer cell in a patient which comprises administering to the patient a composition which reduces integrin-integrin interaction, integrin receptor clustering interaction, or integrin-non-integrin protein interaction.

90. A method of reducing the proliferation of a cancer cell in a patient which comprises administering to the patient a composition which reduces the production of an integrin or laminin by the cancer cell.

91. A method of reducing the proliferation of a cancer cell in a patient which comprises administering to the patient a composition which interferes with an integrin signaling pathway.

92. The methods of claims 88-91, wherein the reduction in cancer cell proliferation is selective.

93. The method of any of claims 88-91, wherein the administering is local or systemic.

94. The method of any of claims 88-91, wherein the patient is additionally administered an agent which modulates cell signaling, a chemotherapeutic drug, or treated with radiation or angiogenesis inhibitor.

95. The method of claim 94, wherein the additional agent is administered serially or in combination.

96. The method of claim 93, wherein the local administering is direct application to cancer cells.

97. The method of claim 96, wherein the direct application to cancer cells is performed during surgery.

98. The method of claim 97, wherein the direct application to cancer cells is performed topically to cancerous tissue.

99. The method of claim 94, wherein the systemic administration is by subcutaneous, intraperitoneal, intra-arterial, intravenous, or bolus administration, or by application through a catheter or similar apparatus.

100. The method of claim 94, wherein the systemic administration comprises a long-term release formulation.

101. The method of claim 94, wherein the systemic administration is by oral administration.

102. The method of claim 101 wherein the oral administration comprises administering a pill, capsule, tablet, liquid or suspension.

103. A method of reducing the proliferation of a cancer cell in a patient which comprises administering to a patient a vector comprising coding sequence for a protein or peptide which inhibits ligand binding to integrin receptors on cancer cells wherein the coding sequence is under the control of a promoter which functions in mammalian cells.

104. A method of reducing the proliferation of a cancer cell in a patient which comprises administering to a patient a vector comprising coding sequence for a protein or peptide which prevents integrin receptor subunits from interacting with one another, prevents integrin receptor clustering interaction, or prevents integrin receptor subunits from interacting with other proteins on cancer cells wherein the coding sequence is under the control of a promoter which functions in mammalian cells.

105. A method of reducing the proliferation of a cancer cell in a patient which comprises administering to a patient a vector comprising coding sequence for a protein or peptide which prevents integrin receptor subunits from interacting with one another, prevents integrin receptor clustering interaction, or prevents integrin receptor subunits from interacting with other proteins in cancer cells wherein the coding sequence is under the control of a promoter which functions in mammalian cells.

106. A method of reducing the proliferation of a cancer cell in a patient which comprises administering to a patient a vector comprising coding sequence for a protein or peptide which interferes with integrin subunit or laminin production wherein the coding sequence is under the control of a promoter which functions in mammalian cells.

107. A method of reducing the proliferation of a cancer cell in a patient which comprises administering to a patient a vector comprising coding sequence for a protein or peptide which interferes with an integrin signaling pathway of cells wherein the coding sequence is under the control of a promoter which functions in mammalian cells.

108. The methods of claims 103-107, wherein the reduction in cancer cell proliferation is selective.

109. The method of claims 103-107 wherein the vector is administered directly to a cancer cell.

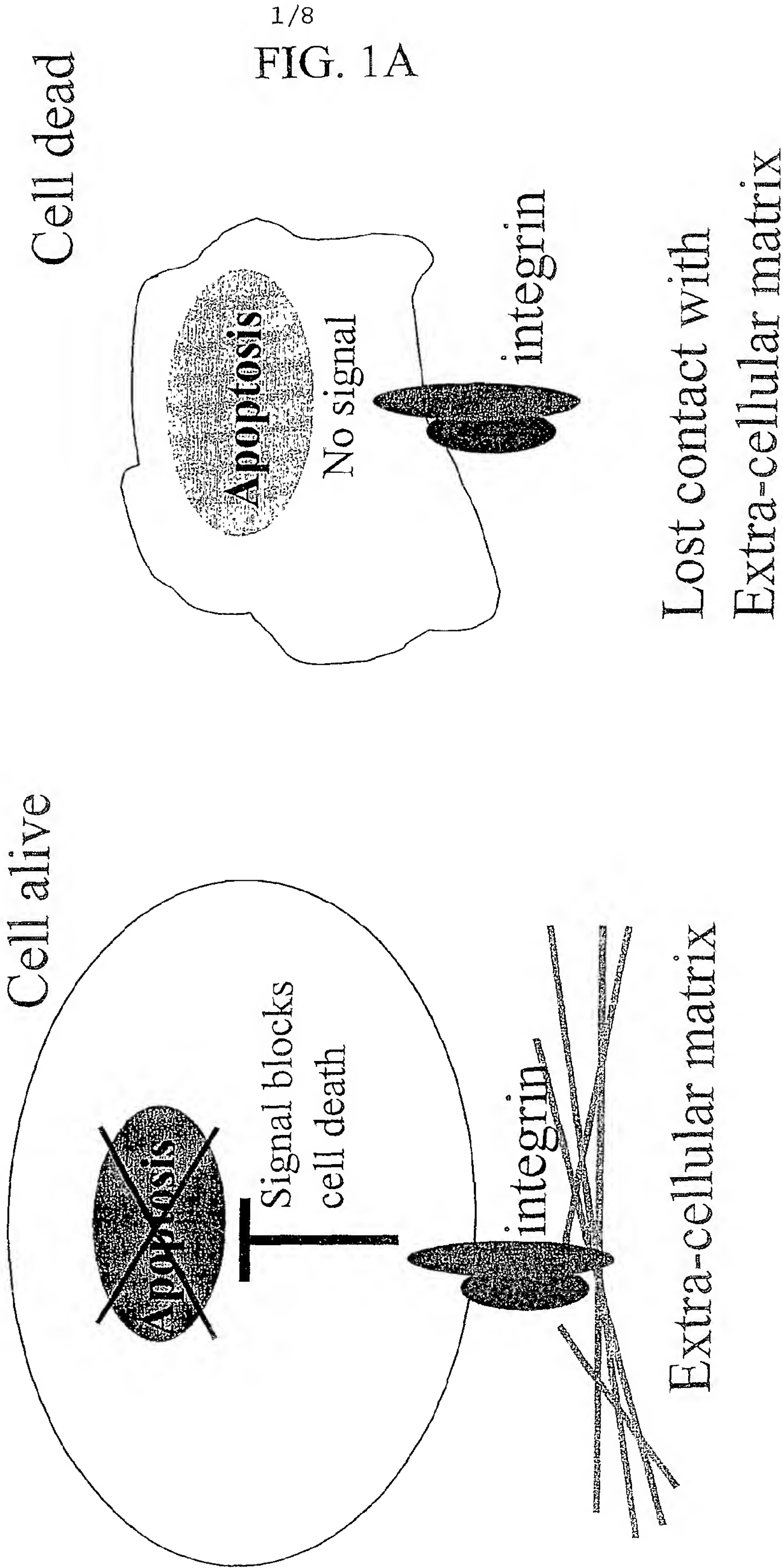
110. The method of claims 103-107, wherein the vector is administered directly to a normal cell.

111. The method of claims 103-107, wherein the vector is packaged in a viral vector or liposome.

112. The method of claim 88 wherein the vector is a retroviral vector.

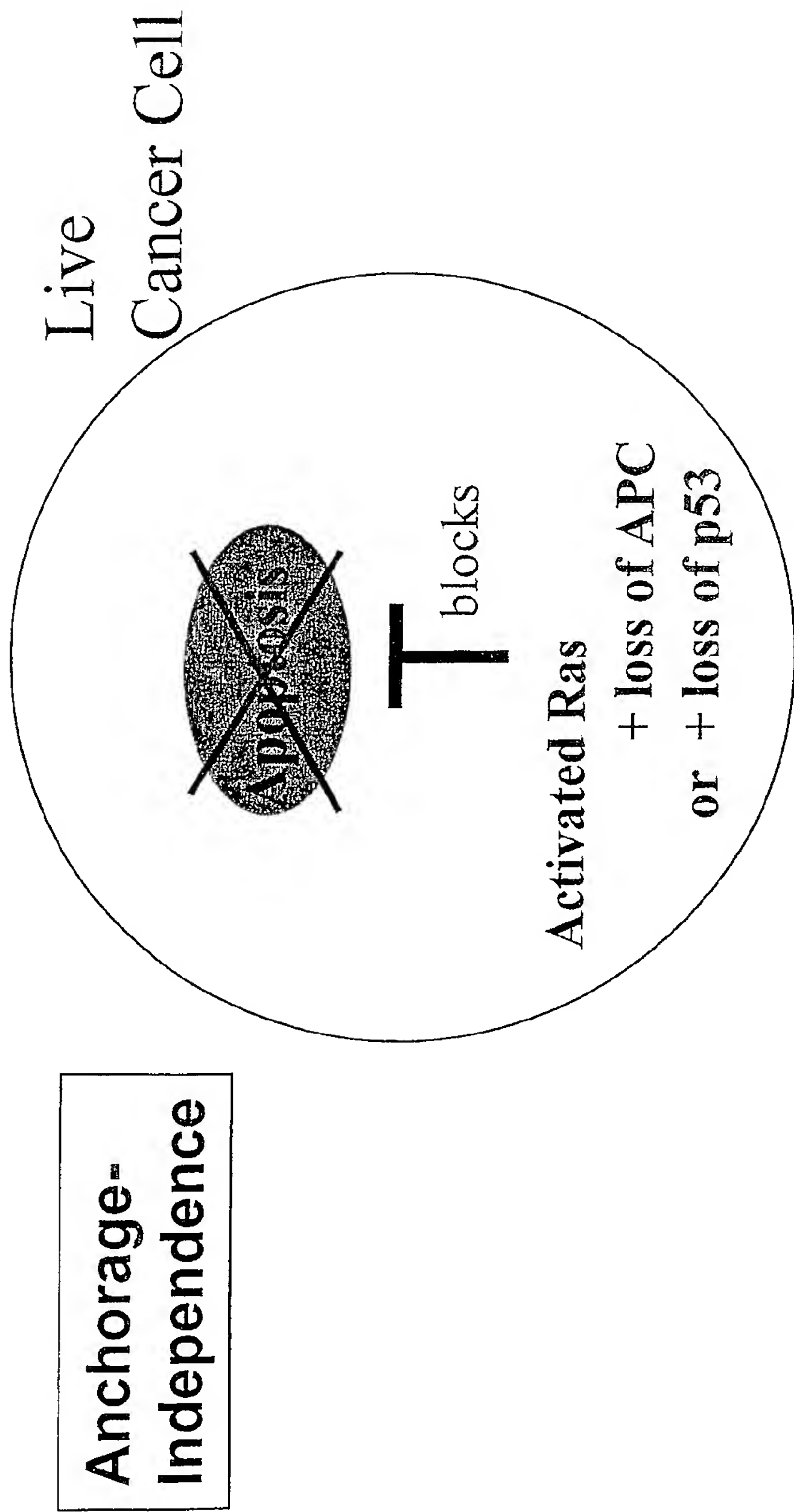
113. The method of any one of claims 103-107, wherein the vector is administered systemically.
114. The method of claim 109, wherein the direct administration is by topical application.
115. The method of claim 111, wherein the direct administration is by topical application.
116. The method of claim 114, wherein the direct administration is performed during surgery.
117. The method of claim 115, wherein the direct administration is performed during surgery.
118. The method of any of claims 103-107, wherein the patient is an animal.
119. The method of claim 118 wherein the animal is a mammal.
120. The method of claim 119, wherein the mammal is human.
121. The method of claims 103-107, wherein the cancer cells are selected from the group consisting of melanoma, adenoma, lymphoma, myeloma, carcinoma, plasmocytoma, sarcoma, glioma, thyoma, leukemia, skin cancer, retinal cancer, breast cancer, prostate cancer, colon cancer, esophageal cancer, stomach cancer, pancreas cancer, brain tumors, lung cancer, ovarian cancer, cervical cancer, hepatic cancer, gastrointestinal cancer, and head and neck cancer cells.
122. The method of any of claims 103-107 wherein the patient is additionally administered at least one of another agent which modulates cell signaling, a chemotherapeutic drug, an angiogenesis inhibitor or treated with radiation.
123. The method of claim 120 wherein the other agent which modifies cell signaling, chemotherapeutic drug, angiogenesis inhibitor or radiation treatment is administered serially or in combination.

Normal cells require integrin-matrix interactions for survival



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FIG. 1A

**Cells transformed by co-operating oncogenic mutations
survive without contact to extra-cellular matrix**



No contact with
Extra-cellular matrix

Panel 2

FIG. 1B

**Oncogenes co-operate to induce cancer cell survival
via engagement of autocrine integrin signaling**

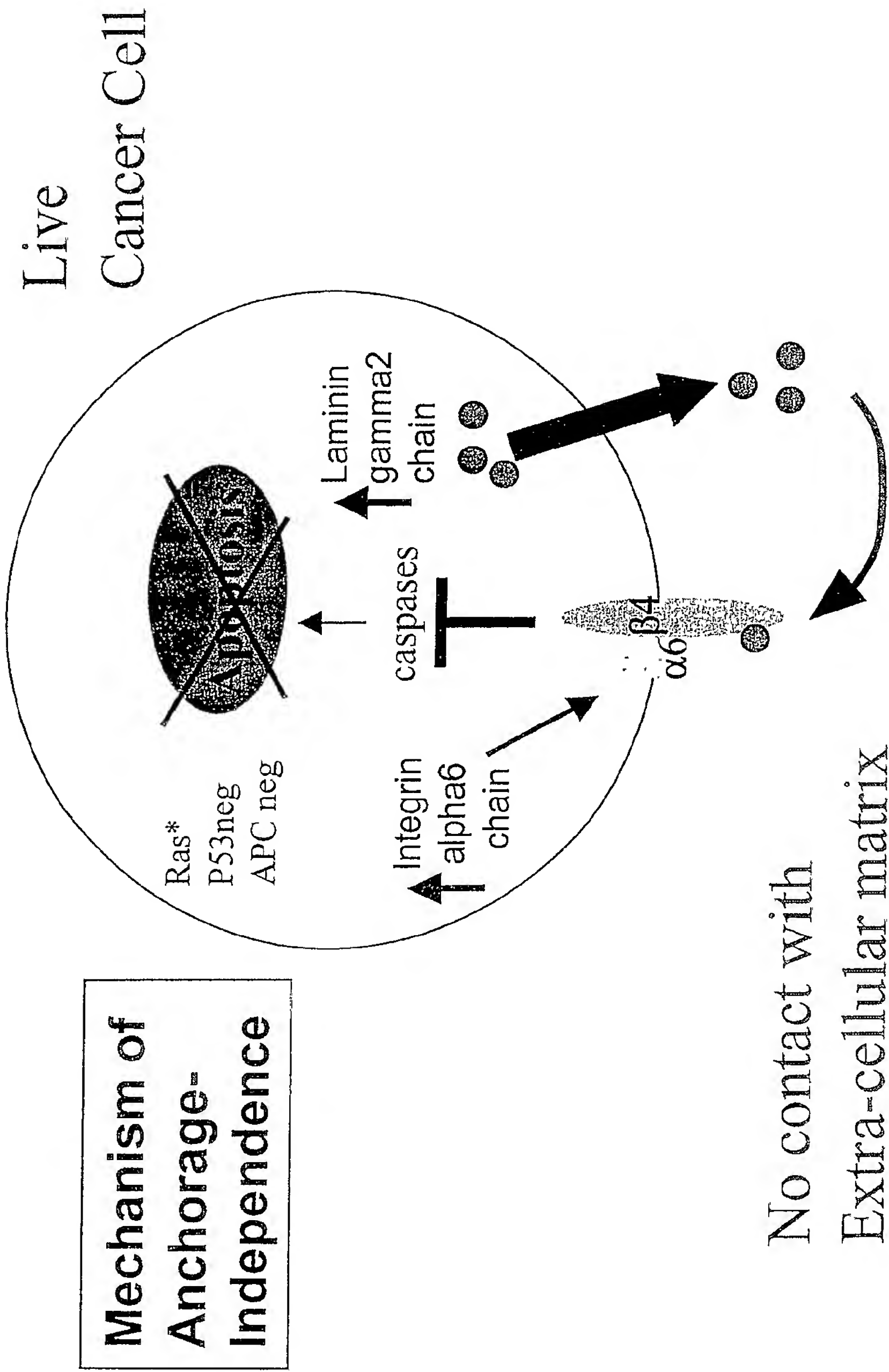


FIG. 1C

Inhibition of oncogene-induced autocrine integrin signaling loop causes cancer cell death

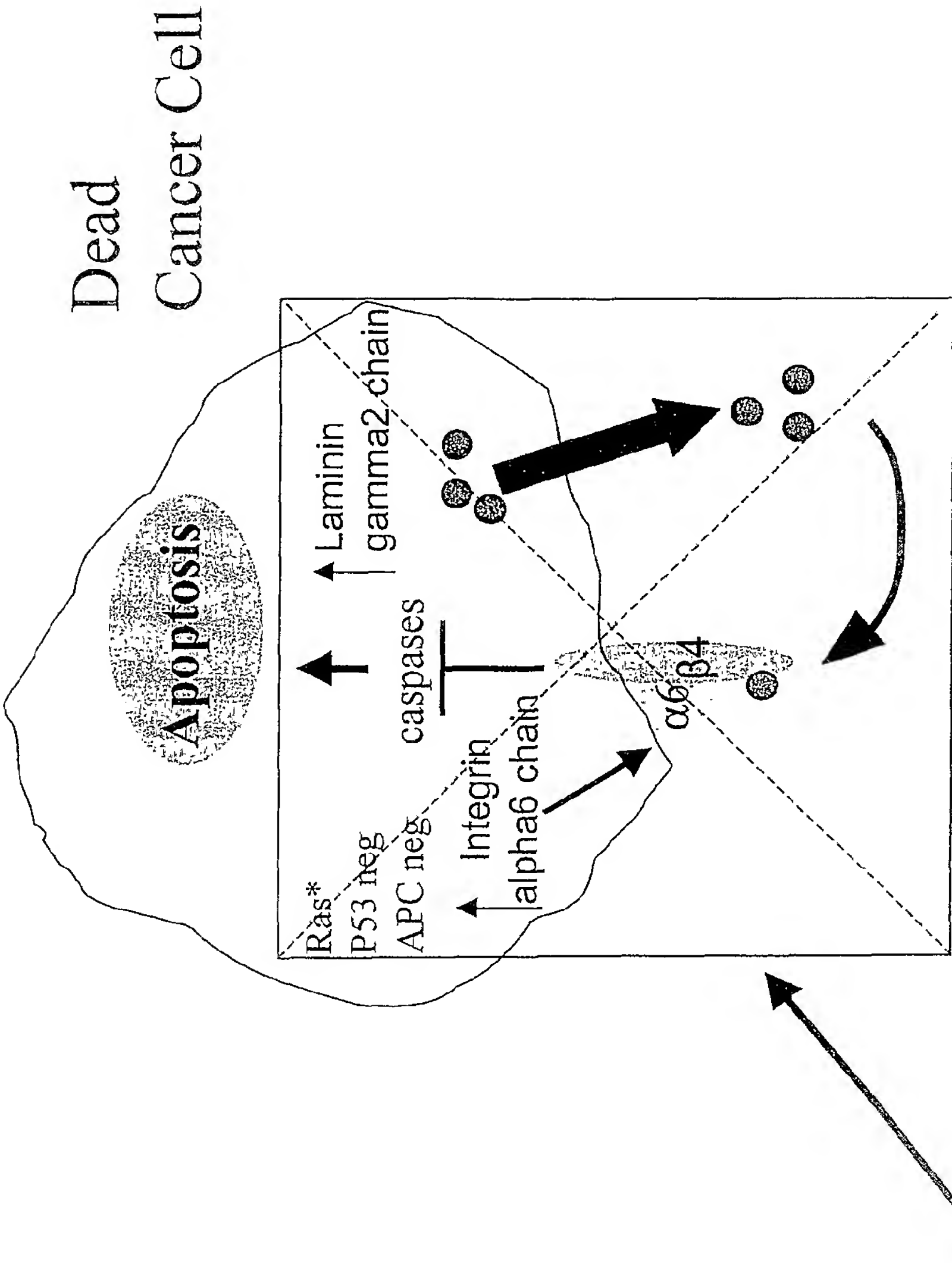


FIG. 1D

Panel 4

5/8
FIG. 2

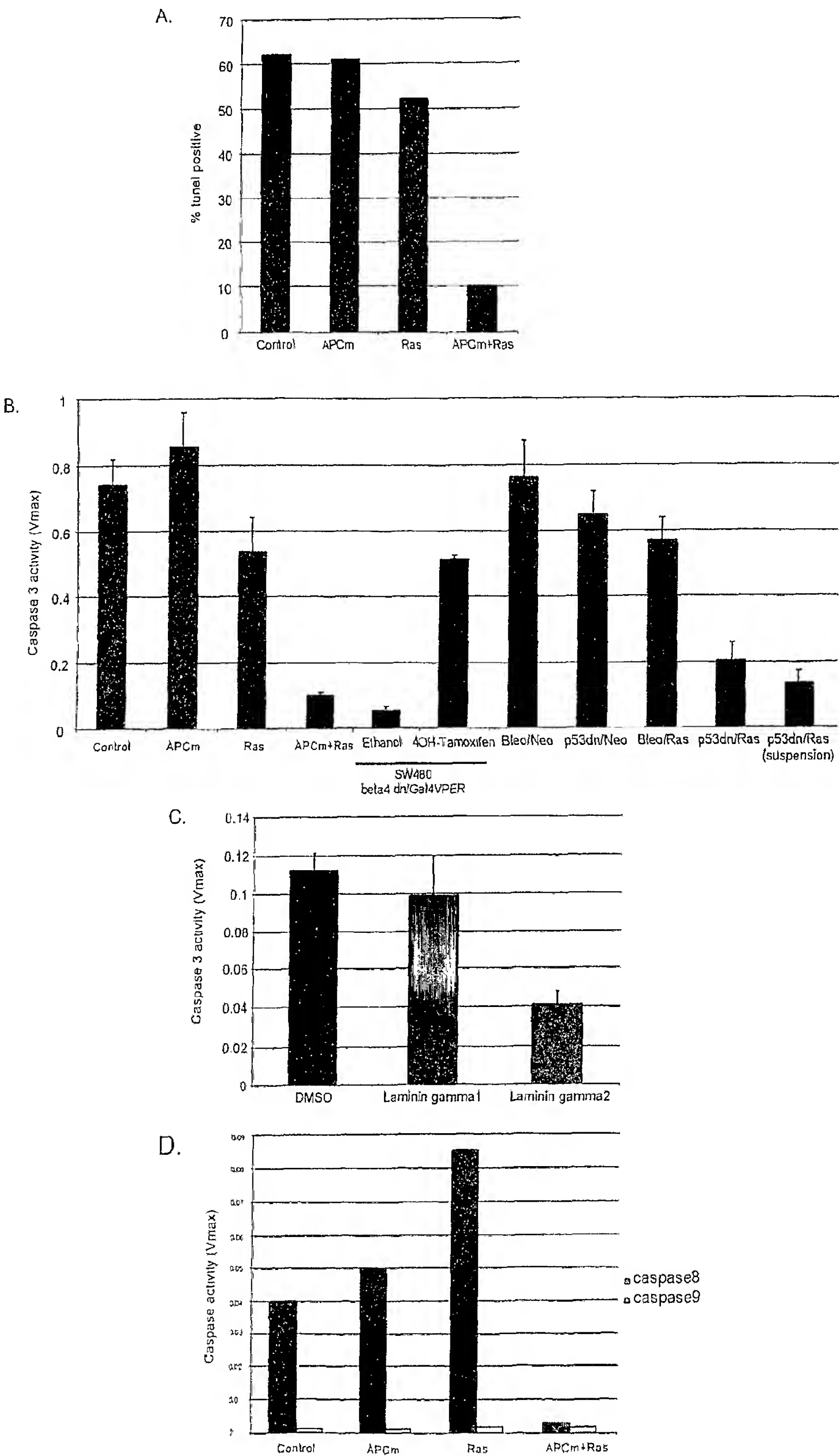


FIG. 3

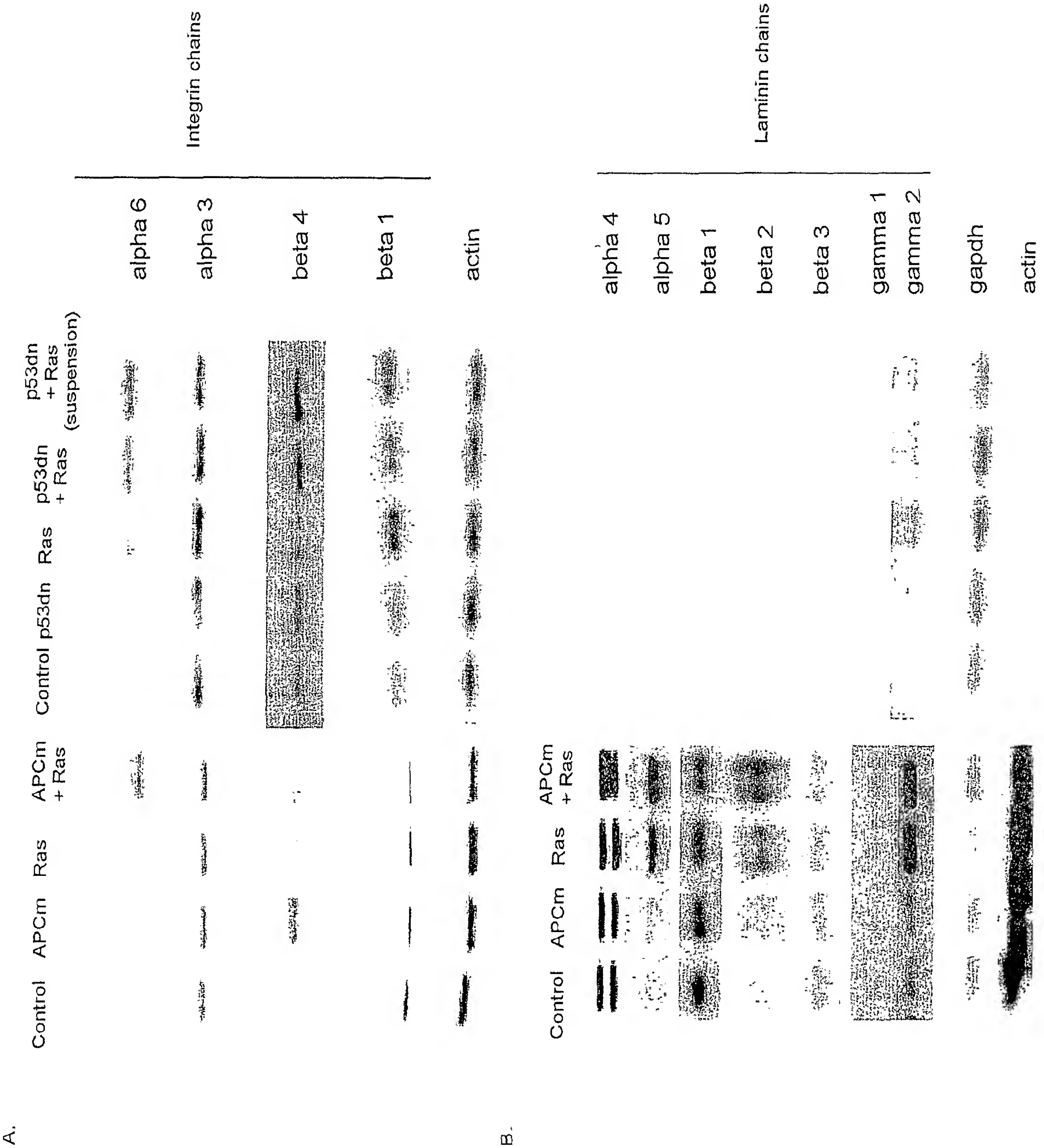


FIG. 4

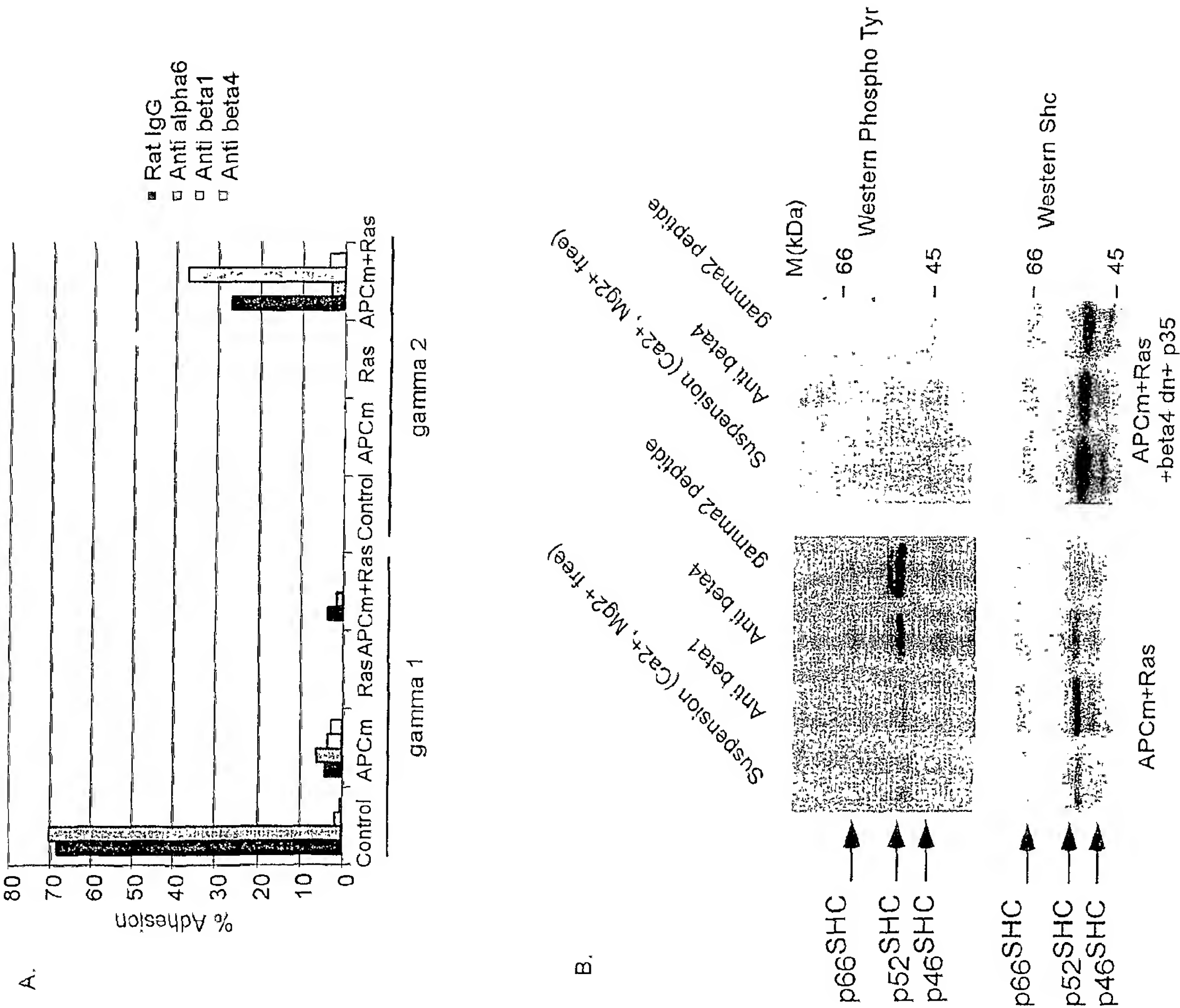
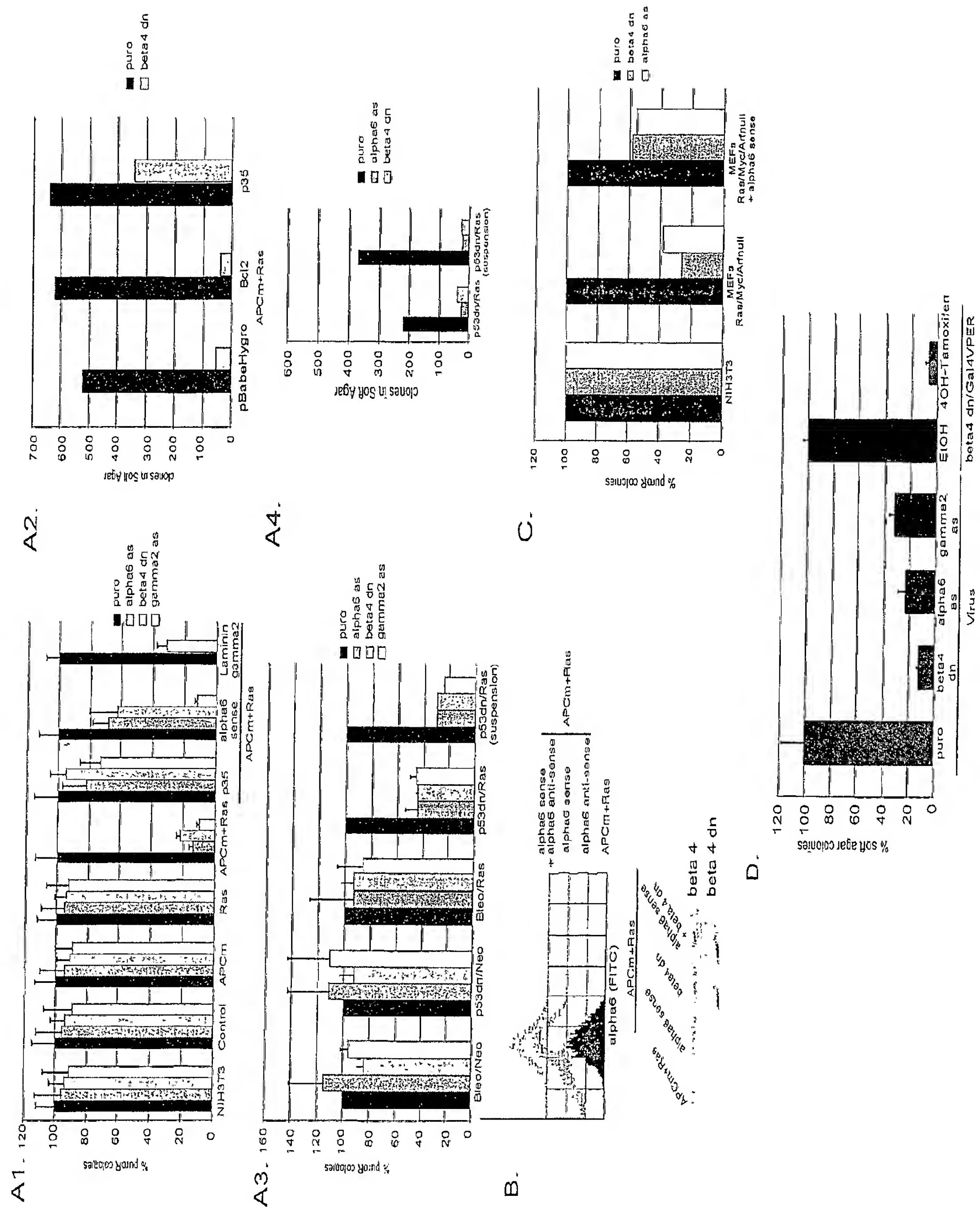


FIG. 5



SEQUENCE LISTING

<110> University of Rochester Medical Center

Deleu, Laurent
Land, Hartmut

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<211> 1073

<212> PRT

<213> Artificial Sequence

<220>

<223> Description of Artificial Sequence; Note =
synthetic construct

<400> 4

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 <223> Description of Artificial Sequence; Note =
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<212> PRT

<213> Artificial Sequence

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12

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<210> 8

<211> 1466

<212> PRT

<213> Artificial Sequence

<220>

<223> Description of Artificial Sequence; Note =
synthetic construct

<400> 8

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Arg Met Leu Ser Ala Ser Leu Pro Gly Asp Leu Ala Asn Arg Cys Lys
          20          25          30
Lys Ala Gln Val Lys Ser Cys Thr Glu Cys Ile Arg Val Asp Lys Ser
          35          40          45
Cys Ala Tyr Cys Thr Asp Glu Leu Phe Lys Glu Arg Arg Cys Asn Thr
          50          55          60

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Gln	Ala	Asp	Val	Leu	Ala	Ala	Gly	Cys	Arg	Gly	Glu	Ser	Ile	Leu	Val
65					70					75					80
Met	Glu	Ser	Ser	Leu	Glu	Ile	Thr	Glu	Asn	Thr	Gln	Ile	Val	Thr	Ser
				85					90					95	
Leu	His	Arg	Ser	Gln	Val	Ser	Pro	Gln	Gly	Leu	Gln	Val	Arg	Leu	Arg
			100					105					110		
Arg	Gly	Glu	Glu	Arg	Thr	Phe	Val	Phe	Gln	Val	Phe	Glu	Pro	Leu	Glu
		115					120					125			
Ser	Pro	Val	Asp	Leu	Tyr	Ile	Leu	Met	Asp	Phe	Ser	Asn	Ser	Met	Ser
	130					135					140				
Asp	Asp	Leu	Asp	Asn	Leu	Lys	Gln	Met	Gly	Gln	Asn	Leu	Ala	Lys	Ile
145				150					155						160
Leu	Arg	Gln	Leu	Thr	Ser	Asp	Tyr	Thr	Ile	Gly	Phe	Gly	Lys	Phe	Val
				165					170					175	
Asp	Lys	Val	Ser	Val	Pro	Gln	Thr	Asp	Met	Arg	Pro	Glu	Lys	Leu	Lys
			180					185					190		
Glu	Pro	Trp	Pro	Asn	Ser	Asp	Pro	Pro	Phe	Ser	Phe	Lys	Asn	Val	Ile
		195					200					205			
Ser	Leu	Thr	Glu	Asn	Val	Glu	Glu	Phe	Trp	Asn	Lys	Leu	Gln	Gly	Glu
	210					215					220				
Arg	Ile	Ser	Gly	Asn	Leu	Asp	Ala	Pro	Glu	Gly	Gly	Phe	Asp	Ala	Ile
225				230						235					240
Leu	Gln	Thr	Ala	Val	Cys	Thr	Arg	Asp	Ile	Gly	Trp	Arg	Ala	Asp	Ser
				245					250					255	
Thr	His	Leu	Leu	Val	Phe	Ser	Thr	Glu	Ser	Ala	Phe	His	Tyr	Glu	Ala
			260					265					270		
Asp	Gly	Ala	Asn	Val	Leu	Ala	Gly	Ile	Met	Asn	Arg	Asn	Asp	Glu	Lys
		275					280					285			
Cys	His	Leu	Asp	Ala	Ser	Gly	Ala	Tyr	Thr	Gln	Tyr	Lys	Thr	Gln	Asp
	290					295					300				
Tyr	Pro	Ser	Val	Pro	Thr	Leu	Val	Arg	Leu	Leu	Ala	Lys	His	Asn	Ile
305					310					315					320
Ile	Pro	Ile	Phe	Ala	Val	Thr	Asn	Tyr	Ser	Tyr	Ser	Tyr	Tyr	Glu	Lys
				325					330					335	
Leu	His	Lys	Tyr	Phe	Pro	Val	Ser	Ser	Leu	Gly	Val	Leu	Gln	Glu	Asp
			340					345					350		
Ser	Ser	Asn	Ile	Val	Glu	Leu	Leu	Glu	Glu	Ala	Phe	Tyr	Arg	Ile	Arg
		355					360					365			
Ser	Asn	Leu	Asp	Ile	Arg	Ala	Leu	Asp	Ser	Pro	Arg	Gly	Leu	Arg	Thr
	370					375					380				
Glu	Val	Thr	Ser	Asp	Thr	Leu	Gln	Lys	Thr	Glu	Thr	Gly	Ser	Phe	His
385					390					395					400
Ile	Lys	Arg	Gly	Glu	Val	Gly	Thr	Tyr	Asn	Val	His	Leu	Arg	Ala	Val
				405					410					415	
Glu	Asp	Ile	Asp	Gly	Thr	His	Val	Cys	Gln	Leu	Ala	Lys	Glu	Asp	Gln
			420					425					430		
Gly	Gly	Asn	Ile	His	Leu	Lys	Pro	Ser	Phe	Ser	Asp	Gly	Leu	Arg	Met
		435					440					445			
Asp	Ala	Ser	Val	Ile	Cys	Asp	Val	Cys	Pro	Cys	Glu	Leu	Gln	Lys	Glu
	450					455					460				
Val	Arg	Ser	Ala	Arg	Cys	His	Phe	Arg	Gly	Asp	Phe	Met	Cys	Gly	His
465					470					475					480
Cys	Val	Cys	Asn	Glu	Gly	Trp	Ser	Gly	Lys	Thr	Cys	Asn	Cys	Ser	Thr
			485						490					495	
Gly	Ser	Leu	Ser	Asp	Thr	Gln	Pro	Cys	Leu	Arg	Glu	Gly	Glu	Asp	Lys
			500					505					510		
Pro	Cys	Ser	Gly	His	Gly	Glu	Cys	Gln	Cys	Gly	Arg	Cys	Val	Cys	Tyr
		515					520					525			
Gly	Glu	Gly	Arg	Tyr	Glu	Gly	His	Phe	Cys	Glu	Tyr	Asp	Asn	Phe	Gln
	530					535					540				
Cys	Pro	Arg	Thr	Ser	Gly	Phe	Leu	Cys	Asn	Asp	Arg	Gly	Arg	Cys	Ser
545					550					555					560

Met	Gly	Glu	Cys	Val	Cys	Glu	Pro	Gly	Trp	Thr	Gly	Arg	Ser	Cys	Asp	
				565					570					575		
Cys	Pro	Leu	Ser	Asn	Ala	Thr	Cys	Ile	Asp	Ser	Asn	Gly	Gly	Ile	Cys	
			580					585					590			
Asn	Gly	Arg	Gly	Tyr	Cys	Glu	Cys	Gly	Arg	Cys	His	Cys	Asn	Gln	Gln	
		595					600					605				
Ser	Leu	Tyr	Thr	Asp	Thr	Thr	Cys	Glu	Ile	Asn	Tyr	Ser	Ala	Ile	Leu	
	610					615					620					
Gly	Leu	Cys	Glu	Asp	Leu	Arg	Ser	Cys	Val	Gln	Cys	Gln	Ala	Trp	Gly	
625					630					635					640	
Thr	Gly	Glu	Lys	Lys	Gly	Arg	Ala	Cys	Asp	Asp	Cys	Pro	Phe	Lys	Val	
			645						650					655		
Lys	Met	Val	Asp	Glu	Leu	Lys	Lys	Glu	Glu	Val	Val	Glu	Tyr	Cys	Ser	
			660					665					670			
Phe	Arg	Asp	Glu	Asp	Asp	Asp	Cys	Thr	Tyr	Ser	Tyr	Asn	Val	Glu	Gly	
		675					680					685				
Asp	Gly	Ser	Pro	Gly	Pro	Asn	Ser	Thr	Val	Leu	Val	His	Lys	Lys	Lys	
	690					695					700					
Asp	Cys	Leu	Pro	Ala	Pro	Ser	Trp	Trp	Leu	Ile	Pro	Leu	Leu	Ile	Phe	
705					710					715					720	
Leu	Leu	Leu	Leu	Leu	Ala	Leu	Leu	Leu	Leu	Leu	Cys	Trp	Lys	Tyr	Cys	
				725					730					735		
Ala	Cys	Cys	Lys	Ala	Cys	Leu	Gly	Leu	Leu	Pro	Cys	Cys	Asn	Arg	Gly	
			740					745					750			
His	Met	Val	Gly	Phe	Lys	Glu	Asp	His	Tyr	Met	Leu	Arg	Glu	Asn	Leu	
		755					760					765				
Met	Ala	Ser	Asp	His	Leu	Asp	Thr	Pro	Met	Leu	Arg	Ser	Gly	Asn	Leu	
	770					775					780					
Lys	Gly	Arg	Asp	Thr	Val	Arg	Trp	Lys	Ile	Thr	Asn	Asn	Val	Gln	Arg	
785					790					795					800	
Pro	Gly	Phe	Ala	Thr	His	Ala	Ala	Ser	Thr	Ser	Pro	Thr	Glu	Leu	Val	
				805					810					815		
Pro	Tyr	Gly	Leu	Ser	Leu	Arg	Leu	Gly	Arg	Leu	Cys	Thr	Glu	Asn	Leu	
			820					825					830			
Met	Lys	Pro	Gly	Thr	Arg	Glu	Cys	Asp	Gln	Leu	Arg	Gln	Glu	Val	Glu	
		835					840					845				
Glu	Asn	Leu	Asn	Glu	Val	Tyr	Arg	Gln	Val	Ser	Gly	Ala	His	Lys	Leu	
	850					855					860					
Gln	Gln	Thr	Lys	Phe	Arg	Gln	Gln	Pro	Asn	Ala	Gly	Lys	Lys	Gln	Asp	
865					870					875					880	
His	Thr	Ile	Val	Asp	Thr	Val	Leu	Leu	Ala	Pro	Arg	Ser	Ala	Lys	Gln	
				885					890					895		
Met	Leu	Leu	Lys	Leu	Thr	Glu	Lys	Gln	Val	Glu	Gln	Gly	Ser	Phe	His	
			900					905					910			
Glu	Leu	Lys	Val	Ala	Pro	Gly	Tyr	Tyr	Thr	Val	Thr	Ala	Glu	Gln	Asp	
		915					920					925				
Ala	Arg	Gly	Met	Val	Glu	Phe	Gln	Glu	Gly	Val	Glu	Leu	Val	Asp	Val	
	930					935					940					
Arg	Val	Pro	Leu	Phe	Ile	Arg	Pro	Glu	Asp	Asp	Asp	Glu	Lys	Gln	Leu	
945					950					955						
Leu	Val	Glu	Ala	Ile	Asp	Val	Pro	Val	Ser	Thr	Ala	Thr	Leu	Gly	Arg	
				965					970					975		
Arg	Leu	Val	Asn	Ile	Thr	Ile	Ile	Lys	Glu	Gln	Ala	Ser	Gly	Val	Val	
			980					985					990			
Ser	Phe	Glu	Gln	Pro	Glu	Tyr	Ser	Val	Ser	Arg	Gly	Asp	Gln	Val	Ala	
		995					1000					1005				
Arg	Ile	Pro	Val	Ile	Arg	His	Ile	Leu	Asp	Asn	Gly	Lys	Ser	Gln	Val	
	1010					1015					1020					
Ser	Tyr	Ser	Thr	Gln	Asp	Asn	Thr	Ala	His	Gly	His	Arg	Asp	Tyr	Val	
1025					1030					1035					1040	
Pro	Val	Glu	Gly	Glu	Leu	Leu	Phe	His	Pro	Gly	Glu	Thr	Trp	Lys	Glu	
				1045					1050					1055		

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Leu Gln Val Lys Leu Leu Glu Leu Gln Glu Val Asp Ser Leu Leu Arg
      1060      1065      1070
Gly Arg Gln Val Arg Arg Phe Gln Val Gln Leu Ser Asn Pro Lys Phe
      1075      1080      1085
Gly Ala Arg Leu Gly Gln Pro Ser Thr Thr Thr Val Ile Leu Asp Glu
      1090      1095      1100
Thr Asp Arg Ser Leu Ile Asn Gln Thr Leu Ser Ser Pro Pro Pro Pro
1105      1110      1115      1120
His Gly Asp Leu Gly Ala Pro Gln Asn Pro Asn Ala Lys Ala Ala Gly
      1125      1130      1135
Ser Arg Lys Ile His Phe Asn Trp Leu Pro Pro Pro Gly Lys Pro Met
      1140      1145      1150
Gly Tyr Arg Val Lys Tyr Trp Ile Gln Gly Asp Ser Glu Ser Glu Ala
      1155      1160      1165
His Leu Leu Asp Ser Lys Val Pro Ser Val Glu Leu Thr Asn Leu Tyr
      1170      1175      1180
Pro Tyr Cys Asp Tyr Glu Met Lys Val Cys Ala Tyr Gly Ala Lys Gly
1185      1190      1195      1200
Glu Gly Pro Tyr Ser Ser Leu Val Ser Cys Arg Thr His Gln Glu Val
      1205      1210      1215
Pro Ser Glu Pro Gly Arg Leu Ala Phe Asn Val Val Ser Ser Thr Val
      1220      1225      1230
Thr Gln Leu Ser Trp Ala Glu Pro Ala Glu Thr Asn Gly Glu Ile Thr
      1235      1240      1245
Ala Tyr Glu Val Cys Tyr Gly Leu Val Asn Glu Asp Asn Arg Pro Ile
      1250      1255      1260
Gly Pro Met Lys Lys Val Leu Val Asp Asn Pro Lys Asn Arg Met Leu
1265      1270      1275      1280
Leu Ile Glu Asn Leu Arg Asp Ser Gln Pro Tyr Arg Tyr Thr Val Lys
      1285      1290      1295
Ala Arg Asn Gly Ala Gly Trp Gly Pro Glu Arg Glu Ala Ile Ile Asn
      1300      1305      1310
Leu Ala Thr Gln Pro Lys Arg Pro Met Ser Ile Pro Ile Ile Pro Asp
      1315      1320      1325
Ile Pro Ile Val Asp Ala Gln Gly Gly Glu Asp Tyr Glu Asn Phe Leu
      1330      1335      1340
Met Tyr Ser Asp Asp Val Leu Arg Ser Pro Ala Ser Ser Gln Arg Pro
1345      1350      1355      1360
Ser Val Ser Asp Asp Thr Glu His Leu Val Asn Gly Arg Met Asp Phe
      1365      1370      1375
Ala Tyr Pro Gly Ser Ala Asn Ser Leu His Arg Met Thr Ala Ala Asn
      1380      1385      1390
Val Ala Tyr Gly Thr His Leu Ser Pro His Leu Ser His Arg Val Leu
      1395      1400      1405
Ser Thr Ser Ser Thr Leu Thr Arg Asp Tyr His Ser Leu Thr Arg Thr
      1410      1415      1420
Glu His Ser His Ser Gly Thr Leu Pro Arg Asp Tyr Ser Thr Leu Thr
1425      1430      1435      1440
Ser Leu Ser Ser Gln Ala Ser Leu Leu Ser Gly Lys Met Gly Gly Ala
      1445      1450      1455
Gly Phe Arg Cys Pro Gly Leu Leu Gly Pro
      1460      1465

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<210> 9

<211> 2205

<212> DNA

<213> Artificial Sequence

<220>

<223> Description of Artificial Sequence; Note =
synthetic construct

<400> 9
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 gaggatcatc ggggtggacaa gagctgtgcc tactgcacag acgagctgtt caaggagagg 180
 cgctgcaaca cccaggcgga cgttctggct gcaggctgca ggggagagag catcctggtc 240
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 caggatcttc cccaaggcct gcaagtccgg ctgaggcggg gtgaggagcg cacgtttgtg 360
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 aactccatgt ctgacgatct ggacaacctc aagcagatgg ggcagaacct ggccaagatc 480
 ctgcccagc tcaccagcga ctacaccatt ggatttgga agtttgtgga caaagtcagc 540
 gtcccacaga cagacatgag gcccgagaaa ctgaaggagc cctggcccaa cagtgatccc 600
 ccgttctcct tcaagaacgt tatcagctta acggagaatg tggaagaatt ctggaacaaa 660
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 ctgcagacag ctgtgtgcac aaggacatt ggctggaggg ctgacagcac ccacctgctg 780
 gtgttctcca ccgagctctg cttccactac gaggctgatg gtgccaacgt tctggccggc 840
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 aagacacagg actaccatc agtgcccacg ctggttcgcc tgcttgccaa gcataacatc 960
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 atcaagcggg gggaggtggg cacatacaat gtgcatctcc gggcagtgga ggacatagat 1260
 gggacacatg tgtgccagct ggctaaagaa gaccaagggg gcaacatcca cctgaaaccc 1320
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 gacacacagc cctgcctgag tgagggtgag gacaaacctg gctcgggcca cggcgagtgc 1560
 cagtgcggac gctgtgtgtg ctatggtgaa ggccgctacg agggtcactt ctgcgagtat 1620
 gacaacttcc agtgtccccg gacctctgga ttctgtgca atgaccgggg acgctgttct 1680
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 caaaaaaga aagactgcct cccggctcct tctgtgtggc tcatccccct gctcatcttc 2160
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<210> 10

<211> 734

<212> PRT

<213> Artificial Sequence

<220>

<223> Description of Artificial Sequence; Note =
 synthetic construct

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			20					25					30		
Lys	Ala	Gln	Val	Lys	Ser	Cys	Thr	Glu	Cys	Ile	Arg	Val	Asp	Lys	Ser
		35					40					45			
Cys	Ala	Tyr	Cys	Thr	Asp	Glu	Leu	Phe	Lys	Glu	Arg	Arg	Cys	Asn	Thr
	50					55					60				
Gln	Ala	Asp	Val	Leu	Ala	Ala	Gly	Cys	Arg	Gly	Glu	Ser	Ile	Leu	Val
65					70					75				80	
Met	Glu	Ser	Ser	Leu	Glu	Ile	Thr	Glu	Asn	Thr	Gln	Ile	Val	Thr	Ser
				85					90					95	

20

Leu	His	Arg	Ser	Gln	Val	Ser	Pro	Gln	Gly	Leu	Gln	Val	Arg	Leu	Arg
			100					105					110		
Arg	Gly	Glu	Glu	Arg	Thr	Phe	Val	Phe	Gln	Val	Phe	Glu	Pro	Leu	Glu
		115					120					125			
Ser	Pro	Val	Asp	Leu	Tyr	Ile	Leu	Met	Asp	Phe	Ser	Asn	Ser	Met	Ser
	130					135					140				
Asp	Asp	Leu	Asp	Asn	Leu	Lys	Gln	Met	Gly	Gln	Asn	Leu	Ala	Lys	Ile
145					150					155					160
Leu	Arg	Gln	Leu	Thr	Ser	Asp	Tyr	Thr	Ile	Gly	Phe	Gly	Lys	Phe	Val
				165					170					175	
Asp	Lys	Val	Ser	Val	Pro	Gln	Thr	Asp	Met	Arg	Pro	Glu	Lys	Leu	Lys
			180					185					190		
Glu	Pro	Trp	Pro	Asn	Ser	Asp	Pro	Pro	Phe	Ser	Phe	Lys	Asn	Val	Ile
		195					200					205			
Ser	Leu	Thr	Glu	Asn	Val	Glu	Glu	Phe	Trp	Asn	Lys	Leu	Gln	Gly	Glu
	210					215					220				
Arg	Ile	Ser	Gly	Asn	Leu	Asp	Ala	Pro	Glu	Gly	Gly	Phe	Asp	Ala	Ile
225					230					235					240
Leu	Gln	Thr	Ala	Val	Cys	Thr	Arg	Asp	Ile	Gly	Trp	Arg	Ala	Asp	Ser
				245					250					255	
Thr	His	Leu	Leu	Val	Phe	Ser	Thr	Glu	Ser	Ala	Phe	His	Tyr	Glu	Ala
			260					265					270		
Asp	Gly	Ala	Asn	Val	Leu	Ala	Gly	Ile	Met	Asn	Arg	Asn	Asp	Glu	Lys
		275					280					285			
Cys	His	Leu	Asp	Ala	Ser	Gly	Ala	Tyr	Thr	Gln	Tyr	Lys	Thr	Gln	Asp
	290					295					300				
Tyr	Pro	Ser	Val	Pro	Thr	Leu	Val	Arg	Leu	Leu	Ala	Lys	His	Asn	Ile
305					310					315					320
Ile	Pro	Ile	Phe	Ala	Val	Thr	Asn	Tyr	Ser	Tyr	Ser	Tyr	Tyr	Glu	Lys
				325					330					335	
Leu	His	Lys	Tyr	Phe	Pro	Val	Ser	Ser	Leu	Gly	Val	Leu	Gln	Glu	Asp
			340					345					350		
Ser	Ser	Asn	Ile	Val	Glu	Leu	Leu	Glu	Glu	Ala	Phe	Tyr	Arg	Ile	Arg
		355					360					365			
Ser	Asn	Leu	Asp	Ile	Arg	Ala	Leu	Asp	Ser	Pro	Arg	Gly	Leu	Arg	Thr
	370					375					380				
Glu	Val	Thr	Ser	Asp	Thr	Leu	Gln	Lys	Thr	Glu	Thr	Gly	Ser	Phe	His
385					390					395					400
Ile	Lys	Arg	Gly	Glu	Val	Gly	Thr	Tyr	Asn	Val	His	Leu	Arg	Ala	Val
				405					410					415	
Glu	Asp	Ile	Asp	Gly	Thr	His	Val	Cys	Gln	Leu	Ala	Lys	Glu	Asp	Gln
			420					425					430		
Gly	Gly	Asn	Ile	His	Leu	Lys	Pro	Ser	Phe	Ser	Asp	Gly	Leu	Arg	Met
		435					440					445			
Asp	Ala	Ser	Val	Ile	Cys	Asp	Val	Cys	Pro	Cys	Glu	Leu	Gln	Lys	Glu
	450					455					460				
Val	Arg	Ser	Ala	Arg	Cys	His	Phe	Arg	Gly	Asp	Phe	Met	Cys	Gly	His
465					470					475					480
Cys	Val	Cys	Asn	Glu	Gly	Trp	Ser	Gly	Lys	Thr	Cys	Asn	Cys	Ser	Thr
				485					490					495	
Gly	Ser	Leu	Ser	Asp	Thr	Gln	Pro	Cys	Leu	Arg	Glu	Gly	Glu	Asp	Lys
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Pro	Cys	Ser	Gly	His	Gly	Glu	Cys	Gln	Cys	Gly	Arg	Cys	Val	Cys	Tyr
		515					520					525			
Gly	Glu	Gly	Arg	Tyr	Glu	Gly	His	Phe	Cys	Glu	Tyr	Asp	Asn	Phe	Gln
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Cys	Pro	Arg	Thr	Ser	Gly	Phe	Leu	Cys	Asn	Asp	Arg	Gly	Arg	Cys	Ser
545					550					555					560
Met	Gly	Glu	Cys	Val	Cys	Glu	Pro	Gly	Trp	Thr	Gly	Arg	Ser	Cys	Asp
				565					570					575	
Cys	Pro	Leu	Ser	Asn	Ala	Thr	Cys	Ile	Asp	Ser	Asn	Gly	Gly	Ile	Cys
			580					585					590		

Asn	Gly	Arg	Gly	Tyr	Cys	Glu	Cys	Gly	Arg	Cys	His	Cys	Asn	Gln	Gln
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Ser	Leu	Tyr	Thr	Asp	Thr	Thr	Cys	Glu	Ile	Asn	Tyr	Ser	Ala	Ile	Leu
	610					615				620					
Gly	Leu	Cys	Glu	Asp	Leu	Arg	Ser	Cys	Val	Gln	Cys	Gln	Ala	Trp	Gly
625					630					635					640
Thr	Gly	Glu	Lys	Lys	Gly	Arg	Ala	Cys	Asp	Asp	Cys	Pro	Phe	Lys	Val
			645					650						655	
Lys	Met	Val	Asp	Glu	Leu	Lys	Lys	Glu	Val	Val	Glu	Tyr	Cys	Ser	
		660					665					670			
Phe	Arg	Asp	Glu	Asp	Asp	Asp	Cys	Thr	Tyr	Ser	Tyr	Asn	Val	Glu	Gly
	675						680					685			
Asp	Gly	Ser	Pro	Gly	Pro	Asn	Ser	Thr	Val	Leu	Val	His	Lys	Lys	Lys
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Asp	Cys	Leu	Pro	Ala	Pro	Ser	Trp	Trp	Leu	Ile	Pro	Leu	Leu	Ile	Phe
705					710					715					720
Leu	Leu	Leu	Leu	Leu	Ala	Leu	Leu	Leu	Leu	Leu	Cys	Trp	Lys		
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<210> 11
 <211> 3579
 <212> DNA
 <213> Artificial Sequence

<220>
 <223> Description of Artificial Sequence; Note =
 synthetic construct

<400> 11

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<210> 12

<211> 1192

<212> PRT

<213> Artificial Sequence

<220>

<223> Description of Artificial Sequence; Note =
synthetic construct

<400> 12

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Lys Ser Arg Gln Cys Val Phe Asp Gln Glu Leu His Arg Gln Ala Gly
 35          40          45
Ser Gly Phe Arg Cys Leu Asn Cys Asn Asp Asn Thr Ala Gly Val His
 50          55          60
Cys Glu Arg Ser Arg Glu Gly Phe Tyr Gln His Gln Ser Lys Ser Arg
 65          70          75          80
Cys Leu Pro Cys Asn Cys His Ser Lys Gly Ser Leu Ser Ala Gly Cys
 85          90          95
Asp Asn Ser Gly Gln Cys Arg Cys Lys Pro Gly Val Thr Gly Gln Arg
100          105          110
Cys Asp Gln Cys Gln Pro Gly Phe His Met Leu Thr Asp Ala Gly Cys
115          120          125
Thr Arg Asp Gln Gly Gln Leu Asp Ser Lys Cys Asp Cys Asp Pro Ala
130          135          140
Gly Ile Ser Gly Pro Cys Asp Ser Gly Arg Cys Val Cys Lys Pro Ala
145          150          155          160
Val Thr Gly Glu Arg Cys Asp Arg Cys Arg Pro Arg Asp Tyr His Leu
165          170          175
Asp Arg Ala Asn Pro Glu Gly Cys Thr Gln Cys Phe Cys Tyr Gly His
180          185          190

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Ser	Gln	Glu	Asn	Asp	Tyr	Lys	Thr	Arg	Leu	Asp	Asp	Leu	Lys	Met	Thr
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Ala	Glu	Arg	Ile	Arg	Ala	Leu	Gly	Ser	Gln	His	Gln	Asn	Arg	Val	Gln
705					710					715					720
Asp	Thr	Ser	Arg	Leu	Ile	Ser	Gln	Met	Arg	Leu	Ser	Leu	Ala	Gly	Ser
				725					730					735	
Glu	Ala	Leu	Leu	Glu	Asn	Thr	Asn	Ile	His	Ser	Ser	Glu	His	Tyr	Val
			740					745					750		
Gly	Pro	Asn	Asp	Phe	Lys	Ser	Leu	Ala	Gln	Glu	Ala	Thr	Arg	Lys	Ala
		755					760					765			
Asp	Ser	His	Ala	Glu	Ser	Ala	Asn	Ala	Met	Lys	Gln	Leu	Ala	Arg	Glu
	770					775					780				
Thr	Glu	Asp	Tyr	Ser	Lys	Gln	Ala	Leu	Ser	Leu	Ala	Arg	Lys	Leu	Leu
785					790					795					800
Ser	Gly	Gly	Gly	Gly	Ser	Gly	Ser	Trp	Asp	Ser	Ser	Val	Val	Gln	Gly
				805					810					815	
Leu	Met	Gly	Lys	Leu	Glu	Lys	Thr	Lys	Ser	Leu	Ser	Gln	Gln	Leu	Ser
			820					825					830		
Leu	Glu	Gly	Thr	Gln	Ala	Asp	Ile	Glu	Ala	Asp	Arg	Ser	Tyr	Gln	His
		835					840					845			
Ser	Leu	Arg	Leu	Leu	Asp	Ser	Ala	Ser	Gln	Leu	Gln	Gly	Val	Ser	Asp
	850					855					860				
Leu	Ser	Phe	Gln	Val	Glu	Ala	Lys	Arg	Ile	Arg	Gln	Lys	Ala	Asp	Ser
865					870					875					880
Leu	Ser	Asn	Leu	Val	Thr	Arg	Gln	Thr	Asp	Ala	Phe	Thr	Arg	Val	Arg
			885						890					895	
Asn	Asn	Leu	Gly	Asn	Trp	Glu	Lys	Glu	Thr	Arg	Gln	Leu	Leu	Gln	Thr
			900					905						910	
Gly	Lys	Asp	Arg	Arg	Gln	Thr	Ser	Asp	Gln	Leu	Leu	Ser	Arg	Ala	Asn
		915					920					925			
Leu	Ala	Lys	Asn	Arg	Ala	Gln	Glu	Ala	Leu	Ser	Met	Gly	Asn	Ala	Thr
	930					935					940				
Phe	Tyr	Glu	Val	Glu	Asn	Ile	Leu	Lys	Asn	Leu	Arg	Glu	Phe	Asp	Leu
945					950					955					960
Gln	Val	Glu	Asp	Arg	Lys	Ala	Glu	Ala	Glu	Glu	Ala	Met	Lys	Arg	Leu
				965					970					975	
Ser	Ser	Ile	Ser	Gln	Lys	Val	Ala	Asp	Ala	Ser	Asp	Lys	Thr	Gln	Gln
			980					985					990		
Ala	Glu	Thr	Ala	Leu	Gly	Ser	Ala	Thr	Ala	Asp	Thr	Gln	Arg	Ala	Lys
		995					1000					1005			
Asn	Ala	Ala	Arg	Glu	Ala	Leu	Glu	Ile	Ser	Ser	Glu	Ile	Glu	Leu	Glu
	1010					1015					1020				
Ile	Gly	Ser	Leu	Asn	Leu	Glu	Ala	Asn	Val	Thr	Ala	Asp	Gly	Ala	Leu
1025					1030					1035					1040
Ala	Met	Glu	Lys	Gly	Thr	Ala	Thr	Leu	Lys	Ser	Glu	Met	Arg	Glu	Met
				1045					1050					1055	
Ile	Glu	Leu	Ala	Arg	Lys	Glu	Leu	Glu	Phe	Asp	Thr	Asp	Lys	Asp	Thr
			1060						1065				1070		
Val	Gln	Leu	Val	Ile	Thr	Glu	Ala	Gln	Gln	Ala	Asp	Ala	Arg	Ala	Thr
		1075					1080					1085			
Ser	Ala	Gly	Val	Thr	Ile	Gln	Asp	Thr	Leu	Asn	Thr	Leu	Asp	Gly	Ile
	1090					1095					1100				
Leu	His	Leu	Ile	Asp	Gln	Pro	Gly	Ser	Val	Asp	Glu	Glu	Gly	Met	Met
1105					1110					1115					1120
Leu	Leu	Glu	Gln	Gly	Leu	Phe	Gln	Ala	Lys	Thr	Gln	Ile	Asn	Ser	Arg
				1125					1130					1135	
Leu	Arg	Pro	Leu	Met	Ser	Asp	Leu	Glu	Glu	Arg	Val	Arg	Arg	Gln	Arg
			1140					1145					1150		
Asn	His	Leu	His	Leu	Leu	Glu	Thr	Ser	Ile	Asp	Gly	Ile	Leu	Ala	Asp
	1155						1160					1165			
Val	Lys	Asn	Leu	Glu	Asn	Ile	Arg	Asp	Asn	Leu	Pro	Pro	Gly	Cys	Tyr
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1185 1190

<210> 13
<211> 3582
<212> DNA
<213> Artificial Sequence

<220>
<223> Description of Artificial Sequence; Note =
synthetic construct

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cgggaacttc acagacaaac tggtaattga ttccgctgcc tcaactgcaa tgacaacact 180
gatggcattc actgcgagaa gtgcaagaat ggcttttacc ggcacagaga aagggaccgc 240
tgtttgccct gcaattgtaa ctccaaaggt tctcttagtg ctcgatgtga caactctgga 300
cgggtgcagct gtaaaccagg tgtgacagga gccagatgog accgatgtct gccaggcttc 360
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<210> 14

<211> 1193

<212> PRT

<213> Artificial Sequence

<220>

<223> Description of Artificial Sequence; Note =
synthetic construct

<400> 14

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 20      25      30
Lys  Ser  Arg  Gln  Cys  Ile  Phe  Asp  Arg  Glu  Leu  His  Arg  Gln  Thr  Gly
 35      40      45
Asn  Gly  Phe  Arg  Cys  Leu  Asn  Cys  Asn  Asp  Asn  Thr  Asp  Gly  Ile  His
 50      55      60
Cys  Glu  Lys  Cys  Lys  Asn  Gly  Phe  Tyr  Arg  His  Arg  Glu  Arg  Asp  Arg
 65      70      75      80
Cys  Leu  Pro  Cys  Asn  Cys  Asn  Ser  Lys  Gly  Ser  Leu  Ser  Ala  Arg  Cys
 85      90      95
Asp  Asn  Ser  Gly  Arg  Cys  Ser  Cys  Lys  Pro  Gly  Val  Thr  Gly  Ala  Arg
100      105      110
Cys  Asp  Arg  Cys  Leu  Pro  Gly  Phe  His  Met  Leu  Thr  Asp  Ala  Gly  Cys
115      120      125
Thr  Gln  Asp  Gln  Arg  Leu  Leu  Asp  Ser  Lys  Cys  Asp  Cys  Asp  Pro  Ala
130      135      140
Gly  Ile  Ala  Gly  Pro  Cys  Asp  Ala  Gly  Arg  Cys  Val  Cys  Lys  Pro  Ala
145      150      155      160
Val  Thr  Gly  Glu  Arg  Cys  Asp  Arg  Cys  Arg  Ser  Gly  Tyr  Tyr  Asn  Leu
165      170      175
Asp  Gly  Gly  Asn  Pro  Glu  Gly  Cys  Thr  Gln  Cys  Phe  Cys  Tyr  Gly  His
180      185      190
Ser  Ala  Ser  Cys  Arg  Ser  Ser  Ala  Glu  Tyr  Ser  Val  His  Lys  Ile  Thr
195      200      205
Ser  Thr  Phe  His  Gln  Asp  Val  Asp  Gly  Trp  Lys  Ala  Val  Gln  Arg  Asn
210      215      220
Gly  Ser  Pro  Ala  Lys  Leu  Gln  Trp  Ser  Gln  Arg  His  Gln  Asp  Val  Phe
225      230      235      240
Ser  Ser  Ala  Gln  Arg  Leu  Asp  Pro  Val  Tyr  Phe  Val  Ala  Pro  Ala  Lys
245      250      255
Phe  Leu  Gly  Asn  Gln  Gln  Val  Ser  Tyr  Gly  Gln  Ser  Leu  Ser  Phe  Asp
260      265      270
Tyr  Arg  Val  Asp  Arg  Gly  Gly  Arg  His  Pro  Ser  Ala  His  Asp  Val  Ile
275      280      285
Leu  Glu  Gly  Ala  Gly  Leu  Arg  Ile  Thr  Ala  Pro  Leu  Met  Pro  Leu  Gly
290      295      300
Lys  Thr  Leu  Pro  Cys  Gly  Leu  Thr  Lys  Thr  Tyr  Thr  Phe  Arg  Leu  Asn
305      310      315      320

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Glu	His	Pro	Ser	Asn	Asn	Trp	Ser	Pro	Gln	Leu	Ser	Tyr	Phe	Glu	Tyr
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Arg	Arg	Leu	Leu	Arg	Asn	Leu	Thr	Ala	Leu	Arg	Ile	Arg	Ala	Thr	Tyr
			340					345					350		
Gly	Glu	Tyr	Ser	Thr	Gly	Tyr	Ile	Asp	Asn	Val	Thr	Leu	Ile	Ser	Ala
		355					360					365			
Arg	Pro	Val	Ser	Gly	Ala	Pro	Ala	Pro	Trp	Val	Glu	Gln	Cys	Ile	Cys
	370					375					380				
Pro	Val	Gly	Tyr	Lys	Gly	Gln	Phe	Cys	Gln	Asp	Cys	Ala	Ser	Gly	Tyr
385					390					395					400
Lys	Arg	Asp	Ser	Ala	Arg	Leu	Gly	Pro	Phe	Gly	Thr	Cys	Ile	Pro	Cys
				405					410					415	
Asn	Cys	Gln	Gly	Gly	Gly	Ala	Cys	Asp	Pro	Asp	Thr	Gly	Asp	Cys	Tyr
			420					425					430		
Ser	Gly	Asp	Glu	Asn	Pro	Asp	Ile	Glu	Cys	Ala	Asp	Cys	Pro	Ile	Gly
		435					440					445			
Phe	Tyr	Asn	Asp	Pro	His	Asp	Pro	Arg	Ser	Cys	Lys	Pro	Cys	Pro	Cys
	450					455					460				
His	Asn	Gly	Phe	Ser	Cys	Ser	Val	Met	Pro	Glu	Thr	Glu	Glu	Val	Val
465					470					475					480
Cys	Asn	Asn	Cys	Pro	Pro	Gly	Val	Thr	Gly	Ala	Arg	Cys	Glu	Leu	Cys
				485					490					495	
Ala	Asp	Gly	Tyr	Phe	Gly	Asp	Pro	Phe	Gly	Glu	His	Gly	Pro	Val	Arg
			500					505					510		
Pro	Cys	Gln	Pro	Cys	Gln	Cys	Asn	Asn	Asn	Val	Asp	Pro	Ser	Ala	Ser
		515					520					525			
Gly	Asn	Cys	Asp	Arg	Leu	Thr	Gly	Arg	Cys	Leu	Lys	Cys	Ile	His	Asn
	530					535					540				
Thr	Ala	Gly	Ile	Tyr	Cys	Asp	Gln	Cys	Lys	Ala	Gly	Tyr	Phe	Gly	Asp
545					550					555					560
Pro	Leu	Ala	Pro	Asn	Pro	Ala	Asp	Lys	Cys	Arg	Ala	Cys	Asn	Cys	Asn
				565					570					575	
Pro	Met	Gly	Ser	Glu	Pro	Val	Gly	Cys	Arg	Ser	Asp	Gly	Thr	Cys	Val
			580					585					590		
Cys	Lys	Pro	Gly	Phe	Gly	Gly	Pro	Asn	Cys	Glu	His	Gly	Ala	Phe	Ser
		595					600					605			
Cys	Pro	Ala	Cys	Tyr	Asn	Gln	Val	Lys	Ile	Gln	Met	Asp	Gln	Phe	Met
	610					615				620					
Gln	Gln	Leu	Gln	Arg	Met	Glu	Ala	Leu	Ile	Ser	Lys	Ala	Gln	Gly	Gly
625					630					635					640
Asp	Gly	Val	Val	Pro	Asp	Thr	Glu	Leu	Glu	Gly	Arg	Met	Gln	Gln	Ala
				645					650					655	
Glu	Gln	Ala	Leu	Gln	Asp	Ile	Leu	Arg	Asp	Ala	Gln	Ile	Ser	Glu	Gly
			660					665					670		
Ala	Ser	Arg	Ser	Leu	Gly	Leu	Gln	Leu	Ala	Lys	Val	Arg	Ser	Gln	Glu
			675				680					685			
Asn	Ser	Tyr	Gln	Ser	Arg	Leu	Asp	Asp	Leu	Lys	Met	Thr	Val	Glu	Arg
	690					695					700				
Val	Arg	Ala	Leu	Gly	Ser	Gln	Tyr	Gln	Asn	Arg	Val	Arg	Asp	Thr	His
705					710					715					720
Arg	Leu	Ile	Thr	Gln	Met	Gln	Leu	Ser	Leu	Ala	Glu	Ser	Glu	Ala	Ser
				725					730					735	
Leu	Gly	Asn	Thr	Asn	Ile	Pro	Ala	Ser	Asp	His	Tyr	Val	Gly	Pro	Asn
			740					745					750		
Gly	Phe	Lys	Ser	Leu	Ala	Gln	Glu	Ala	Thr	Arg	Leu	Ala	Glu	Ser	His
		755					760					765			
Val	Glu	Ser	Ala	Ser	Asn	Met	Glu	Gln	Leu	Thr	Arg	Glu	Thr	Glu	Asp
	770					775					780				
Tyr	Ser	Lys	Gln	Ala	Leu	Ser	Leu	Val	Arg	Lys	Ala	Leu	His	Glu	Gly
785					790					795					800
Val	Gly	Ser	Gly	Ser	Gly	Ser	Pro	Asp	Gly	Ala	Val	Val	Gln	Gly	Leu
				805					310					815	

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Val Glu Lys Leu Glu Lys Thr Lys Ser Leu Ala Gln Gln Leu Thr Arg
      820      825      830
Glu Ala Thr Gln Ala Glu Ile Glu Ala Asp Arg Ser Tyr Gln His Ser
      835      840      845
Leu Arg Leu Leu Asp Ser Val Ser Arg Leu Gln Gly Val Ser Asp Gln
      850      855      860
Ser Phe Gln Val Glu Glu Ala Lys Arg Ile Lys Gln Lys Ala Asp Ser
      865      870      875      880
Leu Ser Thr Leu Val Thr Arg His Met Asp Glu Phe Lys Arg Thr Gln
      885      890      895
Lys Asn Leu Gly Asn Trp Lys Glu Glu Ala Gln Gln Leu Leu Gln Asn
      900      905      910
Gly Lys Ser Gly Arg Glu Lys Ser Asp Gln Leu Leu Ser Arg Ala Asn
      915      920      925
Leu Ala Lys Ser Arg Ala Gln Glu Ala Leu Ser Met Gly Asn Ala Thr
      930      935      940
Phe Tyr Glu Val Glu Ser Ile Leu Lys Asn Leu Arg Glu Phe Asp Leu
      945      950      955      960
Gln Val Asp Asn Arg Lys Ala Glu Ala Glu Glu Ala Met Lys Arg Leu
      965      970      975
Ser Tyr Ile Ser Gln Lys Val Ser Asp Ala Ser Asp Lys Thr Gln Gln
      980      985      990
Ala Glu Arg Ala Leu Gly Ser Ala Ala Ala Asp Ala Gln Arg Ala Lys
      995      1000      1005
Asn Gly Ala Gly Glu Ala Leu Glu Ile Ser Ser Glu Ile Glu Gln Glu
      1010      1015      1020
Ile Gly Ser Leu Asn Leu Glu Ala Asn Val Thr Ala Asp Gly Ala Leu
      1025      1030      1035      1040
Ala Met Glu Lys Gly Leu Ala Ser Leu Lys Ser Glu Met Arg Glu Val
      1045      1050      1055
Glu Gly Glu Leu Glu Arg Lys Glu Leu Glu Phe Asp Thr Asn Met Asp
      1060      1065      1070
Ala Val Gln Met Val Ile Thr Glu Ala Gln Lys Val Asp Thr Arg Ala
      1075      1080      1085
Lys Asn Ala Gly Val Thr Ile Gln Asp Thr Leu Asn Thr Leu Asp Gly
      1090      1095      1100
Leu Leu His Leu Met Asp Gln Pro Leu Ser Val Asp Glu Glu Gly Leu
      1105      1110      1115      1120
Val Leu Leu Glu Gln Lys Leu Ser Arg Ala Lys Thr Gln Ile Asn Ser
      1125      1130      1135
Gln Leu Arg Pro Met Met Ser Glu Leu Glu Glu Arg Ala Arg Gln Gln
      1140      1145      1150
Arg Gly His Leu His Leu Leu Glu Thr Ser Ile Asp Gly Ile Leu Ala
      1155      1160      1165
Asp Val Lys Asn Leu Glu Asn Ile Arg Asp Asn Leu Pro Pro Gly Cys
      1170      1175      1180
Tyr Asn Thr Gln Ala Leu Glu Gln Gln
      1185      1190

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<210> 15

<211> 1111

<212> PRT

<213> Artificial Sequence

<220>

<223> Description of Artificial Sequence; Note =
synthetic construct

<400> 15

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```


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Lys	Ser	Arg	Gln	Cys	Ile	Phe	Asp	Arg	Glu	Leu	His	Arg	Gln	Thr	Gly
		35					40					45			
Asn	Gly	Phe	Arg	Cys	Leu	Asn	Cys	Asn	Asp	Asn	Thr	Asp	Gly	Ile	His
	50					55					60				
Cys	Glu	Lys	Cys	Lys	Asn	Gly	Phe	Tyr	Arg	His	Arg	Glu	Arg	Asp	Arg
65					70					75					80
Cys	Leu	Pro	Cys	Asn	Cys	Asn	Ser	Lys	Gly	Ser	Leu	Ser	Ala	Arg	Cys
				85					90					95	
Asp	Asn	Ser	Gly	Arg	Cys	Ser	Cys	Lys	Pro	Gly	Val	Thr	Gly	Ala	Arg
			100					105					110		
Cys	Asp	Arg	Cys	Leu	Pro	Gly	Phe	His	Met	Leu	Thr	Asp	Ala	Gly	Cys
		115					120					125			
Thr	Gln	Asp	Gln	Arg	Leu	Leu	Asp	Ser	Lys	Cys	Asp	Cys	Asp	Pro	Ala
						135						140			
Gly	Ile	Ala	Gly	Pro	Cys	Asp	Ala	Gly	Arg	Cys	Val	Cys	Lys	Pro	Ala
145					150					155					160
Val	Thr	Gly	Glu	Arg	Cys	Asp	Arg	Cys	Arg	Ser	Gly	Tyr	Tyr	Asn	Leu
				165					170					175	
Asp	Gly	Gly	Asn	Pro	Glu	Gly	Cys	Thr	Gln	Cys	Phe	Cys	Tyr	Gly	His
			180					185					190		
Ser	Ala	Ser	Cys	Arg	Ser	Ser	Ala	Glu	Tyr	Ser	Val	His	Lys	Ile	Thr
		195					200					205			
Ser	Thr	Phe	His	Gln	Asp	Val	Asp	Gly	Trp	Lys	Ala	Val	Gln	Arg	Asn
		210				215					220				
Gly	Ser	Pro	Ala	Lys	Leu	Gln	Trp	Ser	Gln	Arg	His	Gln	Asp	Val	Phe
225					230					235					240
Ser	Ser	Ala	Gln	Arg	Leu	Asp	Pro	Val	Tyr	Phe	Val	Ala	Pro	Ala	Lys
				245					250					255	
Phe	Leu	Gly	Asn	Gln	Gln	Val	Ser	Tyr	Gly	Gln	Ser	Leu	Ser	Phe	Asp
			260					265					270		
Tyr	Arg	Val	Asp	Arg	Gly	Gly	Arg	His	Pro	Ser	Ala	His	Asp	Val	Ile
		275					280					285			
Leu	Glu	Gly	Ala	Gly	Leu	Arg	Ile	Thr	Ala	Pro	Leu	Met	Pro	Leu	Gly
	290					295					300				
Lys	Thr	Leu	Pro	Cys	Gly	Leu	Thr	Lys	Thr	Tyr	Thr	Phe	Arg	Leu	Asn
305					310					315					320
Glu	His	Pro	Ser	Asn	Asn	Trp	Ser	Pro	Gln	Leu	Ser	Tyr	Phe	Glu	Tyr
				325					330					335	
Arg	Arg	Leu	Leu	Arg	Asn	Leu	Thr	Ala	Leu	Arg	Ile	Arg	Ala	Thr	Tyr
			340					345					350		
Gly	Glu	Tyr	Ser	Thr	Gly	Tyr	Ile	Asp	Asn	Val	Thr	Leu	Ile	Ser	Ala
		355					360					365			
Arg	Pro	Val	Ser	Gly	Ala	Pro	Ala	Pro	Trp	Val	Glu	Gln	Cys	Ile	Cys
		370				375					380				
Pro	Val	Gly	Tyr	Lys	Gly	Gln	Phe	Cys	Gln	Asp	Cys	Ala	Ser	Gly	Tyr
385					390					395					400
Lys	Arg	Asp	Ser	Ala	Arg	Leu	Gly	Pro	Phe	Gly	Thr	Cys	Ile	Pro	Cys
				405					410					415	
Asn	Cys	Gln	Gly	Gly	Gly	Ala	Cys	Asp	Pro	Asp	Thr	Gly	Asp	Cys	Tyr
			420					425					430		
Ser	Gly	Asp	Glu	Asn	Pro	Asp	Ile	Glu	Cys	Ala	Asp	Cys	Pro	Ile	Gly
		435					440					445			
Phe	Tyr	Asn	Asp	Pro	His	Asp	Pro	Arg	Ser	Cys	Lys	Pro	Cys	Pro	Cys
	450					455					460				
His	Asn	Gly	Phe	Ser	Cys	Ser	Val	Met	Pro	Glu	Thr	Glu	Glu	Val	Val
465					470					475					480
Cys	Asn	Asn	Cys	Pro	Pro	Gly	Val	Thr	Gly	Ala	Arg	Cys	Glu	Leu	Cys
				485					490					495	
Ala	Asp	Gly	Tyr	Phe	Gly	Asp	Pro	Phe	Gly	Glu	His	Gly	Pro	Val	Arg
			500					505					510		

Pro	Cys	Gln	Pro	Cys	Gln	Cys	Asn	Asn	Asn	Val	Asp	Pro	Ser	Ala	Ser	
		515					520					525				
Gly	Asn	Cys	Asp	Arg	Leu	Thr	Gly	Arg	Cys	Leu	Lys	Cys	Ile	His	Asn	
	530					535					540					
Thr	Ala	Gly	Ile	Tyr	Cys	Asp	Gln	Cys	Lys	Ala	Gly	Tyr	Phe	Gly	Asp	
545					550					555					560	
Pro	Leu	Ala	Pro	Asn	Pro	Ala	Asp	Lys	Cys	Arg	Ala	Cys	Asn	Cys	Asn	
				565					570						575	
Pro	Met	Gly	Ser	Glu	Pro	Val	Gly	Cys	Arg	Ser	Asp	Gly	Thr	Cys	Val	
			580					585					590			
Cys	Lys	Pro	Gly	Phe	Gly	Gly	Pro	Asn	Cys	Glu	His	Gly	Ala	Phe	Ser	
		595					600					605				
Cys	Pro	Ala	Cys	Tyr	Asn	Gln	Val	Lys	Ile	Gln	Met	Asp	Gln	Phe	Met	
	610					615					620					
Gln	Gln	Leu	Gln	Arg	Met	Glu	Ala	Leu	Ile	Ser	Lys	Ala	Gln	Gly	Gly	
625					630					635					640	
Asp	Gly	Val	Val	Pro	Asp	Thr	Glu	Leu	Glu	Gly	Arg	Met	Gln	Gln	Ala	
				645					650						655	
Glu	Gln	Ala	Leu	Gln	Asp	Ile	Leu	Arg	Asp	Ala	Gln	Ile	Ser	Glu	Gly	
			660				665						670			
Ala	Ser	Arg	Ser	Leu	Gly	Leu	Gln	Leu	Ala	Lys	Val	Arg	Ser	Gln	Glu	
		675					680					685				
Asn	Ser	Tyr	Gln	Ser	Arg	Leu	Asp	Asp	Leu	Lys	Met	Thr	Val	Glu	Arg	
	690					695					700					
Val	Arg	Ala	Leu	Gly	Ser	Gln	Tyr	Gln	Asn	Arg	Val	Arg	Asp	Thr	His	
705					710					715					720	
Arg	Leu	Ile	Thr	Gln	Met	Gln	Leu	Ser	Leu	Ala	Glu	Ser	Glu	Ala	Ser	
				725					730						735	
Leu	Gly	Asn	Thr	Asn	Ile	Pro	Ala	Ser	Asp	His	Tyr	Val	Gly	Pro	Asn	
			740				745						750			
Gly	Phe	Lys	Ser	Leu	Ala	Gln	Glu	Ala	Thr	Arg	Leu	Ala	Glu	Ser	His	
		755					760					765				
Val	Glu	Ser	Ala	Ser	Asn	Met	Glu	Gln	Leu	Thr	Arg	Glu	Thr	Glu	Asp	
	770					775					780					
Tyr	Ser	Lys	Gln	Ala	Leu	Ser	Leu	Val	Arg	Lys	Ala	Leu	His	Glu	Gly	
785					790					795					800	
Val	Gly	Ser	Gly	Ser	Gly	Ser	Pro	Asp	Gly	Ala	Val	Val	Gln	Gly	Leu	
				805					810					815		
Val	Glu	Lys	Leu	Glu	Lys	Thr	Lys	Ser	Leu	Ala	Gln	Gln	Leu	Thr	Arg	
			820				825						830			
Glu	Ala	Thr	Gln	Ala	Glu	Ile	Glu	Ala	Asp	Arg	Ser	Tyr	Gln	His	Ser	
		835					840					845				
Leu	Arg	Leu	Leu	Asp	Ser	Val	Ser	Arg	Leu	Gln	Gly	Val	Ser	Asp	Gln	
	850					855					860					
Ser	Phe	Gln	Val	Glu	Glu	Ala	Lys	Arg	Ile	Lys	Gln	Lys	Ala	Asp	Ser	
865					870					875					880	
Leu	Ser	Thr	Leu	Val	Thr	Arg	His	Met	Asp	Glu	Phe	Lys	Arg	Thr	Gln	
				885					890					895		
Lys	Asn	Leu	Gly	Asn	Trp	Lys	Glu	Glu	Ala	Gln	Gln	Leu	Leu	Gln	Asn	
			900					905						910		
Gly	Lys	Ser	Gly	Arg	Glu	Lys	Ser	Asp	Gln	Leu	Leu	Ser	Arg	Ala	Asn	
		915					920					925				
Leu	Ala	Lys	Ser	Arg	Ala	Gln	Glu	Ala	Leu	Ser	Met	Gly	Asn	Ala	Thr	
	930					935					940					
Phe	Tyr	Glu	Val	Glu	Ser	Ile	Leu	Lys	Asn	Leu	Arg	Glu	Phe	Asp	Leu	
945					950					955					960	
Gln	Val	Asp	Asn	Arg	Lys	Ala	Glu	Ala	Glu	Glu	Ala	Met	Lys	Arg	Leu	
				965					970					975		
Ser	Tyr	Ile	Ser	Gln	Lys	Val	Ser	Asp	Ala	Ser	Asp	Lys	Thr	Gln	Gln	
			980					985					990			
Ala	Glu	Arg	Ala	Leu	Gly	Ser	Ala	Ala	Ala	Asp	Ala	Gln	Arg	Ala	Lys	
		995					1000					1005				

Asn Gly Ala Gly Glu Ala Leu Glu Ile Ser Ser Glu Ile Glu Gln Glu
 1010 1015 1020
 Ile Gly Ser Leu Asn Leu Glu Ala Asn Val Thr Ala Asp Gly Ala Leu
 1025 1030 1035 1040
 Ala Met Glu Lys Gly Leu Ala Ser Leu Lys Ser Glu Met Arg Glu Val
 1045 1050 1055
 Glu Gly Glu Leu Glu Arg Lys Glu Leu Glu Phe Asp Thr Asn Met Asp
 1060 1065 1070
 Ala Val Gln Met Val Ile Thr Glu Ala Gln Lys Val Asp Thr Arg Ala
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 Leu Leu His Leu Met Gly Met
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 <213> Artificial Sequence

<220>
 <223> Description of Artificial Sequence; Note =
 synthetic construct

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 aacacccagg cggagctgct ggccgoggggc tgccagcggg agagcatcgt ggtcatggag 240
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 agaagaaggg actgccctcc gggctccttc tgggtggctca tccccctgct cctcctcttc 2160
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<210> 17

<211> 1873

<220>

<223> Description of Artificial Sequence; Note =
synthetic construct

<400> 17

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Pro	Ile	Lys 35	Ser	Cys	Thr	Glu	Cys 40	Val	Arg	Val	Asp	Lys 45	Asp	Cys	Ala
Tyr	Cys 50	Thr	Asp	Glu	Met	Phe 55	Arg	Asp	Arg	Arg	Cys 60	Asn	Thr	Gln	Ala
Glu 65	Leu	Leu	Ala	Ala	Gly 70	Cys	Gln	Arg	Glu	Ser 75	Ile	Val	Val	Met	Glu 80
Ser	Ser	Phe	Gln	Ile 85	Thr	Glu	Glu	Thr	Gln 90	Ile	Asp	Thr	Thr	Leu 95	Arg
Arg	Ser	Gln	Met 100	Ser	Pro	Gln	Gly	Leu 105	Arg	Val	Arg	Leu	Arg 110	Pro	Gly
Glu	Glu	Arg 115	His	Phe	Glu	Leu	Glu 120	Val	Phe	Glu	Pro	Leu 125	Glu	Ser	Pro
Val	Asp 130	Leu	Tyr	Ile	Leu	Met 135	Asp	Phe	Ser	Asn	Ser 140	Met	Ser	Asp	Asp
Leu 145	Asp	Asn	Leu	Lys	Lys 150	Met	Gly	Gln	Asn	Leu 155	Ala	Arg	Val	Leu	Ser 160
Gln	Leu	Thr	Ser	Asp 165	Tyr	Thr	Ile	Gly	Phe 170	Gly	Lys	Phe	Val	Asp 175	Lys
Val	Ser	Val	Pro 180	Gln	Thr	Asp	Met	Arg 185	Pro	Glu	Lys	Leu	Lys 190	Glu	Pro
Trp	Pro	Asn 195	Ser	Asp	Pro	Pro	Phe 200	Ser	Phe	Lys	Asn	Val 205	Ile	Ser	Leu
Thr	Glu 210	Asp	Val	Asp	Glu	Phe 215	Arg	Asn	Lys	Leu	Gln 220	Gly	Glu	Arg	Ile
Ser 225	Gly	Asn	Leu	Asp	Ala 230	Pro	Glu	Gly	Gly	Phe 235	Asp	Ala	Ile	Leu	Gln 240
Thr	Ala	Val	Cys	Thr 245	Arg	Asp	Ile	Gly	Trp 250	Arg	Pro	Asp	Ser	Thr 255	His
Leu	Leu	Val	Phe 260	Ser	Thr	Glu	Ser	Ala 265	Phe	His	Tyr	Glu	Ala 270	Asp	Gly
Ala	Asn 275	Val	Leu	Ala	Gly	Ile	Met 280	Ser	Arg	Asn	Asp	Glu 285	Arg	Cys	His
Leu	Asp 290	Thr	Thr	Gly	Thr	Tyr 295	Thr	Gln	Tyr	Arg	Thr 300	Gln	Asp	Tyr	Pro
Ser 305	Val	Pro	Thr	Leu	Val 310	Arg	Leu	Leu	Ala	Lys 315	His	Asn	Ile	Ile	Pro 320
Ile	Phe	Ala	Val	Thr 325	Asn	Tyr	Ser	Tyr	Ser 330	Tyr	Tyr	Glu	Lys	Leu 335	His
Thr	Tyr	Phe	Pro 340	Val	Ser	Ser	Leu	Gly 345	Val	Leu	Gln	Glu	Asp 350	Ser	Ser
Asn	Ile 355	Val	Glu	Leu	Leu	Glu	Glu 360	Ala	Phe	Asn	Arg	Ile 365	Arg	Ser	Asn
Leu	Asp 370	Ile	Arg	Ala	Leu	Asp 375	Ser	Pro	Arg	Gly	Leu 380	Arg	Thr	Glu	Val
Thr 385	Ser	Lys	Met	Phe	Gln 390	Lys	Thr	Arg	Thr	Gly 395	Ser	Phe	His	Ile	Arg 400
Arg	Gly	Glu	Val	Gly 405	Ile	Tyr	Gln	Val	Gln 410	Leu	Arg	Ala	Leu	Glu 415	His
Val	Asp	Gly	Thr 420	His	Val	Cys	Gln	Leu 425	Pro	Glu	Asp	Gln	Lys 430	Gly	Asn

Ile	His	Leu	Lys	Pro	Ser	Phe	Ser	Asp	Gly	Leu	Lys	Met	Asp	Ala	Gly
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Ile	Ile	Cys	Asp	Val	Cys	Thr	Cys	Glu	Leu	Gln	Lys	Glu	Val	Arg	Ser
	450					455					460				
Ala	Arg	Cys	Ser	Phe	Asn	Gly	Asp	Phe	Val	Cys	Gly	Gln	Cys	Val	Cys
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Ser	Glu	Gly	Trp	Ser	Gly	Gln	Thr	Cys	Asn	Cys	Ser	Thr	Gly	Ser	Leu
				485					490					495	
Ser	Asp	Ile	Gln	Pro	Cys	Leu	Arg	Glu	Gly	Glu	Asp	Lys	Pro	Cys	Ser
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Gly	Arg	Gly	Glu	Cys	Gln	Cys	Gly	His	Cys	Val	Cys	Tyr	Gly	Glu	Gly
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Arg	Tyr	Glu	Gly	Gln	Phe	Cys	Glu	Tyr	Asp	Asn	Phe	Gln	Cys	Pro	Arg
	530					535					540				
Thr	Ser	Gly	Phe	Leu	Cys	Asn	Asp	Arg	Gly	Arg	Cys	Ser	Met	Gly	Gln
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Cys	Val	Cys	Glu	Pro	Gly	Trp	Thr	Gly	Pro	Ser	Cys	Asp	Cys	Pro	Leu
				565					570					575	
Ser	Asn	Ala	Thr	Cys	Ile	Asp	Ser	Asn	Gly	Gly	Ile	Cys	Asn	Gly	Arg
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Gly	His	Cys	Glu	Cys	Gly	Arg	Cys	His	Cys	His	Gln	Gln	Ser	Leu	Tyr
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Thr	Asp	Thr	Ile	Cys	Glu	Ile	Asn	Tyr	Ser	Ala	Ser	Thr	Arg	Ala	Ser
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Trp	Thr	Ser	Leu	Arg	Glu	Ala	Arg	Arg	Trp	Trp	Cys	Ala	Ala	Pro	Ser
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Ala	Pro	Leu	Gly	Pro	Thr	Ala	Leu	Ser	Trp	Cys	Thr	Arg	Arg	Arg	Asp
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Cys	Pro	Pro	Gly	Ser	Phe	Trp	Trp	Leu	Ile	Pro	Leu	Leu	Leu	Leu	Leu
705					710					715					720
Leu	Pro	Leu	Leu	Ala	Leu	Leu	Leu	Leu	Leu	Cys	Trp	Lys	Tyr	Cys	Ala
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Cys	Cys	Lys	Ala	Cys	Leu	Ala	Leu	Leu	Pro	Cys	Cys	Asn	Arg	Gly	His
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Met	Val	Gly	Phe	Lys	Glu	Asp	His	Tyr	Met	Leu	Arg	Glu	Asn	Leu	Met
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Ala	Ser	Asp	His	Leu	Asp	Thr	Pro	Met	Leu	Arg	Ser	Gly	Asn	Leu	Lys
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Gly	Phe	Ala	Thr	His	Ala	Ala	Ser	Ile	Asn	Pro	Thr	Glu	Leu	Val	Pro
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Lys	Pro	Asp	Thr	Arg	Glu	Cys	Ala	Gln	Leu	Arg	Gln	Glu	Val	Glu	Glu
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865					870					875					880
Thr	Ile	Val	Asp	Thr	Val	Leu	Met	Ala	Pro	Arg	Ser	Ala	Lys	Pro	Ala
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Leu	Leu	Lys	Leu	Thr	Glu	Lys	Gln	Val	Glu	Gln	Arg	Ala	Phe	His	Asp
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		915					920					925			

Arg	Gly	Met	Val	Glu	Phe	Gln	Glu	Gly	Val	Glu	Leu	Val	Asp	Val	Arg
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Val	Glu	Ala	Ile	Asp	Val	Pro	Ala	Gly	Thr	Ala	Thr	Leu	Gly	Arg	Arg
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Pro	Glu	Leu	Ile	Pro	Arg	Leu	Ser	Ala	Ser	Ser	Gly	Arg	Ser	Ser	Asp
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 Leu Val Asn Gly Arg Met Asp Phe Ala Phe Pro Gly Ser Thr Asn Ser
 1445 1450 1455
 Leu His Arg Met Thr Thr Thr Ser Ala Ala Ala Tyr Gly Thr His Leu
 1460 1465 1470
 Ser Pro His Val Pro His Arg Val Leu Ser Thr Ser Ser Thr Leu Thr
 1475 1480 1485
 Arg Asp Tyr Asn Ser Leu Thr Arg Ser Glu His Ser His Ser Thr Thr
 1490 1495 1500
 Leu Pro Arg Asp Tyr Ser Thr Leu Thr Ser Val Ser Ser His Gly Leu
 1505 1510 1515 1520
 Pro Pro Ile Trp Glu His Gly Arg Ser Arg Leu Pro Leu Ser Trp Ala
 1525 1530 1535
 Leu Gly Ser Arg Ser Arg Ala Gln Met Lys Gly Phe Pro Pro Ser Arg
 1540 1545 1550
 Gly Pro Arg Asp Ser Ile Ile Leu Ala Gly Arg Pro Ala Ala Pro Ser
 1555 1560 1565
 Trp Gly Pro Asp Ser Arg Leu Thr Ala Gly Val Pro Asp Thr Pro Thr
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 Arg Leu Val Phe Ser Ala Leu Gly Pro Thr Ser Leu Arg Val Ser Trp
 1585 1590 1595 1600
 Gln Glu Pro Arg Cys Glu Arg Pro Leu Gln Gly Tyr Ser Val Glu Tyr
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 Glu Gly Val Ile Thr Ile Glu Ser Gln Val His Pro Gln Ser Pro Leu
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 Cys Pro Leu Pro Gly Ser Ala Phe Thr Leu Ser Thr Pro Ser Ala Pro
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 Gly Pro Leu Val Phe Thr Ala Leu Ser Pro Asp Ser Leu Gln Leu Ser
 1700 1705 1710
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 Thr Cys Glu Met Ala Gln Gly Gly Gly Pro Ala Thr Ala Phe Arg Val
 1730 1735 1740
 Asp Gly Asp Ser Pro Glu Ser Arg Leu Thr Val Pro Gly Leu Ser Glu
 1745 1750 1755 1760
 Asn Val Pro Tyr Lys Phe Lys Val Gln Ala Arg Thr Thr Glu Gly Phe
 1765 1770 1775
 Gly Pro Glu Arg Glu Gly Ile Ile Thr Ile Glu Ser Gln Asp Gly Gly
 1780 1785 1790
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 Gln Ser Glu Tyr Ser Ser Ile Ser Thr Thr His Thr Ser Ala Thr Glu
 1810 1815 1820
 Pro Phe Leu Val Gly Pro Thr Leu Gly Ala Gln His Leu Glu Ala Gly
 1825 1830 1835 1840
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 Thr

<210> 18
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 <212> DNA

<213> Artificial Sequence

<220>

<223> Description of Artificial Sequence; Note =
synthetic construct

<400> 18

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